R-585-4-9-16

SITE INSPECTION OF WASHINGTON PLATING PREPARED UNDER

TDD NO. F3-8810-15 EPA NO. DC-007 CONTRACT NO. 68-01-7346

FOR THE

HAZARDOUS SITE CONTROL DIVISION
U.S. ENVIRONMENTAL PROTECTION AGENCY

SEPTEMBER 13, 1989

NUS CORPORATION SUPERFUND DIVISION

SUBMITTED BY

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REVIEWED BY

(b) (4)

SECTION SUPERVISOR

APPROVED BY

GARTH GLENN

REGIONAL OPERATIONS

MANAGER, FIT 3



Site Name: Washington Plating TDD No.: F3-8810-15

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Site Name: Washington Plating
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(Red)

APPENDICES

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PLATES

1 FOUR-MILE-RADIUS MAP

ORIGINAL (Red)

SECTION 1

TDD No.: <u>F3-8810-15</u>

ORIGINAL (Red)

1.0 INTRODUCTION

1.1 Authorization

NUS Corporation performed this work under Environmental Protection Agency Contract No. 68-01-7346. This specific report was prepared in accordance with Technical Directive Document No. F3-8810-15 for the Washington Plating site, located in Washington, D.C.

1.2 Scope of Work

NUS FIT 3 was tasked to conduct a site inspection of the subject site.

1.3 Summary

The Washington Plating site, located in Washington, D.C., is an inactive, one-acre electroplating facility where automobile bumpers were straightened and refinished. Operations at the site began in 1973 and ended in January 1987.

The nickel and chromium electroplating process utilized 12 tanks with various solutions. Effluent from four rinse tanks was drained into the city sewage system. In 1986, blockage in the sewer line that is under the alley and behind the electroplating room caused rinse effluent to overflow from a manhole. The effluent flowed along the alley, behind neighboring backyards, and ultimately into a storm drain on the corner of 13th Street and V Street. Officials from the District of Columbia Department of Commerce and Regulatory Affairs witnessed one overflow incident on March 10, 1986. Complaints had been filed by a resident on V Street on previous occasions during the preceding months, but no evidence of effluent in the alleys was seen during follow-up investigations.

The rinse tanks were used after the bumpers were dipped into either a nickel- or chromium-plating tank. The tanks were drained approximately two to three times each day.

According to Milan Milosevic, the manager of the facility, no other solutions were drained into the sewer from the plating room. A strip tank using caustic soda with a reverse current was used to strip the bumpers in the straightening room. This tank was drained into the sewer system once every couple of years. It was last drained in February 1986, prior to the relocation of the entire operation. The operation moved to 2215 Adams Place Northeast in Washington, D.C.



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All residents within the site area obtain their drinking water from public sources. The nearest public source is located over 13 miles away. The nearest residence is located 20 feet from the site.

NUS FIT 3 conducted a site inspection on December 6, 1988. Activities included sampling on-site surface water and sediment and off-site soils. The results of sampling, as shown in section 7.0 and discussed in section 8.0, have revealed elevated levels of heavy metals including lead, chromium, and polyaromatic hydrocarbons (PAHs) in on-site sediment samples.

SECTION 2

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ORIGINAL (Red)

2.0 THE SITE

2.1 Location

The site is located in the northwestern section of Washington, D.C., along 14th Street (see figure 2.1, page 2-2). The site can be located on the United States Geological Survey (U.S.G.S.) Washington West,

D.C. - Maryland - Virginia 7.5 minute series quadrangle map at north 38° 55′ 04″ latitude and west 77°

01' 52" longitude, or 7-7/8 inches north and 4-1/2 inches west of the southeastern corner of the map.1

2.2 Site Layout

The site is approximately one acre in size. It slopes gradually (less than one percent) toward the

southeast. The site, which is divided by two alleys, supports three buildings. The site lies on the

eastern side of the 2100 block of 14th Street and is bounded by alleys to the north, east, and south.

Fourteenth Street runs north-south (see figure 2.2, page 2-3).1,2

Between 14th Street and the western alley is a building measuring approximately 135 by 50 feet. This

was the old plating room. Its street address is 2109 14th Street. This building consists of a single open

room with a three-feet-high platform along the north wall. A single drain is located in the center of

the floor. This drain leads to the sewer. Beneath the platform is a basement measuring

approximately 65 by 10 feet. At the time of the site visit, the basement was flooded with

approximately four feet of water.2,3

Across the western alley and west of the eastern alley are two other, larger facility buildings. Both

buildings measure approximately 256 by 56 feet and run east-west. They are separated by a center

alley that is approximately 20 feet wide. The northern building, which is bounded to the north by the

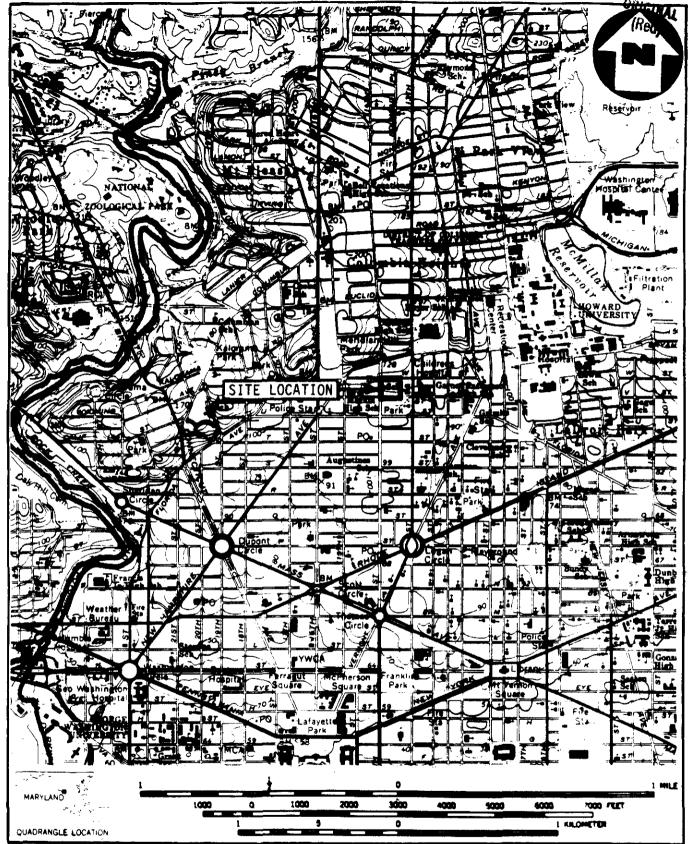
northern alley, was the old straightening and finishing building. This building is currently empty. The

southern building, which is bounded to the south by the southern alley, was the old bumper storage

building. This building was damaged from a fire that occurred in 1988. This building is currently used

for truck storage.^{2,4}

2-1



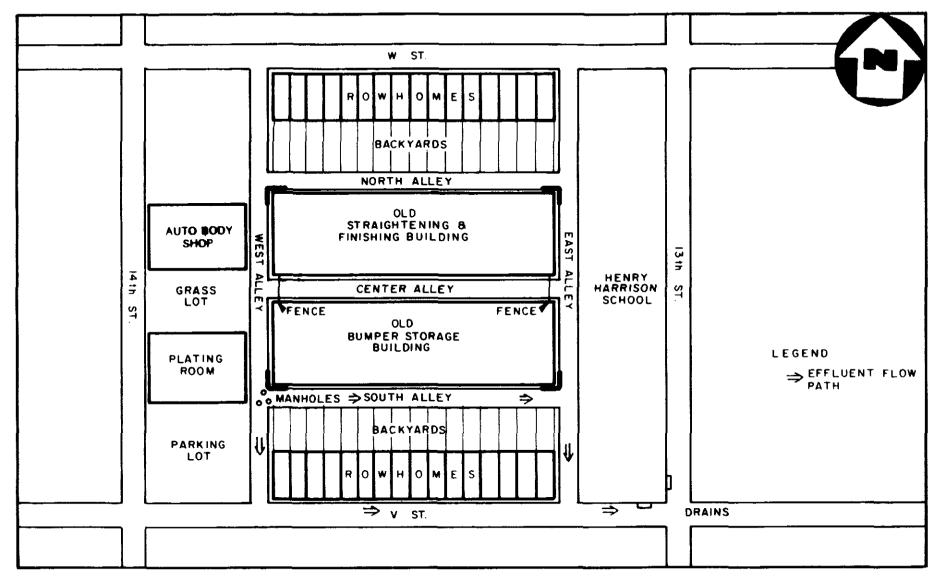
SOURCE: (7.5 MINUTE SERIES) USGS WASHINGTON WEST, D.C. - MD. - VA. QUAD.

SITE LOCATION MAP

WASHINGTON PLATING CO., WASHINGTON, D.C.

SCALE 1:24000





SITE SKETCH

WASHINGTON PLATING CO., WASHINGTON, D.C.
(NO SCALE)





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ORIGINAL (Red)

A sewage line runs along the western alley behind the plating room to the main under V Street. Southeast of the plating room are three manholes that lead to the sewer line.^{2,3}

South of the southern alley and north of the northern alley are row homes. They front on V and W Streets, respectively. The Henry Harrison School is located immediately east of the eastern alley, fronting on 13th Street. South of the plating room is a parking lot. North of the plating room are a grass lot and an auto body shop. 1,2

2.3 Ownership History

George Galich has owned and operated the facility since February 1973. The property had been owned by a laundry and dry cleaning company since 1938.^{5,6}

2.4 Site Use History

George Galich has operated an automobile bumper straightening and replating company since 1973. In January 1986, he moved his straightening and finishing operations to a location in northwest Washington, D.C. The electroplating operations continued at the site until January 1987, when they were also moved to the new location.^{3,5}

The sewer system was used on a daily basis for the direct discharge of rinse water. The four rinse tanks were drained two to three times a day. No pretreatment was performed by the Washington Plating Company. At the time of the FIT 3 preliminary assessment, it was discovered that the sewer connection from the plating room to the sewer line under the alley was blocked. According to Mr. Milosevic, this blockage occurred in September or October 1986. Therefore, during the last two to three months of operations in the plating room, the rinse water had flooded the basement.³

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2.5 <u>Permit and Regulatory Action History</u>

The facility obtained a waste discharge permit from the Department of Public Works for the discharge of its wastewater into the sewer system. Mr. Milosevic tested the effluent in order to file a baseline report with EPA in October 1985. Alex Slinsky, of the Water Management Division, Water Permits Branch, District of Columbia, Maryland, and Virginia section of EPA Region 3, received the report. Mr. Slinsky reported to District of Columbia's Department of Consumer and Regulatory Affairs that Washington Plating Company was within acceptable limits for the categorical limits of pretreatment standards for electroplaters, as set forth in the Federal Register.⁷

In February 1986, the District of Columbia Department of Consumer and Regulatory Affairs received a complaint from a resident on V Street. The report was that a greenish liquid was flowing down a back alley from the Washington Plating Company. The situation was inspected the same day by Byron Bacon, of the department's Waste Management Branch, but no effluent was observed. The report occurred during a period of heavy rain. Subsequently, the government of the District of Columbia sampled soil and liquids in and around the site (see appendix C).

The same resident filed another complaint in March 1986. When Mr. Bacon arrived, the Hazardous Materials Team of the District of Columbia police and fire departments were already at the scene. The Hazardous Materials Team identified the effluent as nickel sulfite. Two samples were taken from puddles in the alley. Mr. Milosevic indicated that, because of a blocked sewer line, the effluent overflowed from a manhole in the alley. The blocked sewer was referred to the Department of Public Works. Subsequently, the government of the District of Columbia sampled soils around the plating facility (see appendix C).⁷

2.6 Remedial Action to Date

In January 1987, the electroplating process and all the supplies and equipment were moved to a new location. No other remedial action has taken place to date.³

SECTION 3

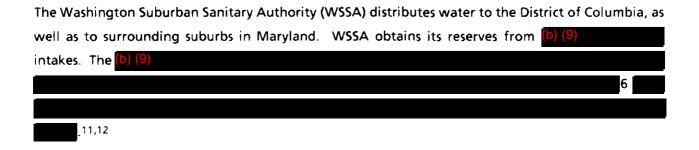
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3.0 ENVIRONMENTAL SETTING

3.1 Water Supply

All residents within the three-mile-radius study area surrounding the Washington Plating site utilize public supplies to obtain potable drinking water.



A small portion of Arlington County, Virginia lies in the southwestern portion of the study area. This area is serviced by the Fairfax County Water Authority (FCWA), which also obtains its water from a surface water intake. This intake is (b) (9)

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The water

volume contributed by these wells constitutes less than 0.5 percent of the total supply obtained from the surface intake. These wells, located outside the 3-mile radius, are drilled to various depths, ranging between 300 and 500 feet beneath the surface, and tap either the Coastal Plain deposits or the Wissahickon Formation. No known privately owned domestic groundwater wells are currently used to obtain potable supplies. However, the average depth of wells across the study area is 123 feet. 12,13,14

3.2 Surface Waters

The storm drainage inlet that collects runoff from the site is located on the corner of 13th. Street and V. Street. The storm drainage system is a combined system with the municipal sewage system. The drainage is treated before discharging into the Potomac River. 9,10



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The Department of Public Works operates an extensive sewage treatment plant along the Potomac River. The water intake for the Washington Suburban Sanitary Commission is where Interstate 70 meets the Potomac, upstream and outside the three-mile radius of the study area.¹⁰

The McMillan Reservoir is located 3/4 mile northeast of the site. The Potomac River is approximately 2-1/2 miles south-southwest of the site. The river is used for recreational purposes, mainly for an access waterway to the Chesapeake Bay. Rock Creek is approximately one mile west of the site.¹

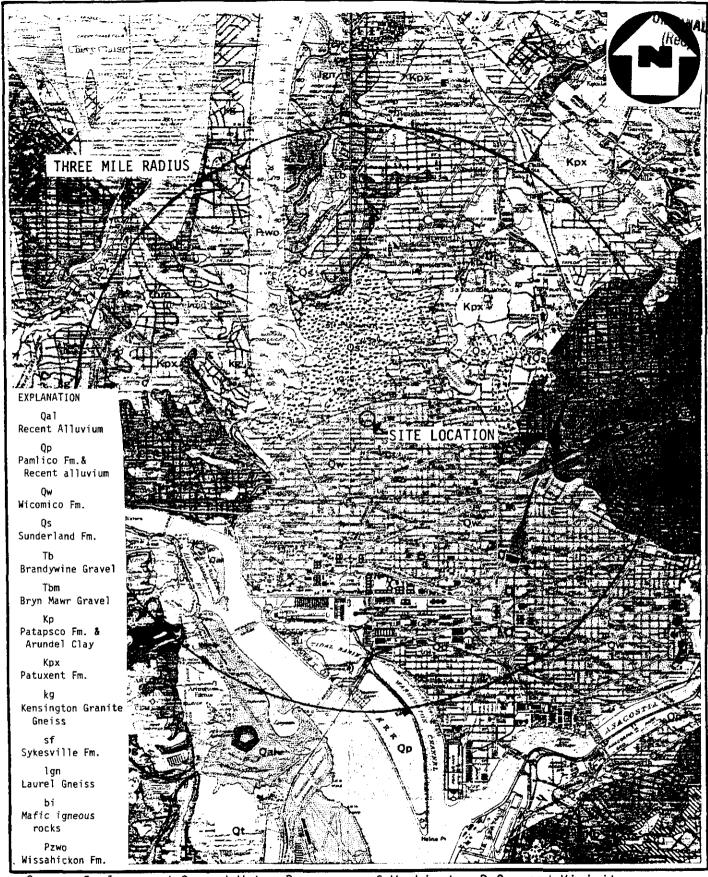
3.3 Hydrogeology

The geologic and hydrogeologic conditions in the study area were researched as part of the site inspection. A preliminary literature review was conducted to determine surface and subsurface geologic conditions, soil character, and the status of groundwater transport and storage.

3.3.1 Geology

The Washington Plating site lies on the Fall Line, the boundary separating the Piedmont Physiographic Province to the northwest from the Coastal Plain Physiographic Province to the southeast. The Piedmont, which is typically composed of hard igneous and metamorphic rocks, underlies and forms the basement complex for the Coastal Plain sediments east of the Fall Line. The major structural feature of the Piedmont region is the schistocity that is commonly present in subsurface lithologies and sub-parallels the regional northeast-southwestwardly trend. No additional structural features such as folding and/or faulting are present within the study area. The unconsolidated sediments of the Coastal Plain form a wedge that gently dips and thickens to the southeast. The ages of the Coastal Plain sediments range from the Cretaceous at the base upward through the Quaternary. The Washington area displays an undulating topography with moderate relief. Elevations across the study area range from sea level to approximately 400 feet above sea level.15

The site is underlain by the Quaternary age Wicomico Formation, consisting of an unconsolidated coarse gravel basal bed, with finer sand and yellow to white silt above. Local deposits of carbonaceous clay that contains woody debris are also present (see figure 3.1, page 3-3). The stratigraphic thickness of the Wicomico rarely exceeds 30 feet.¹⁵



Source:Geology and Ground-Water Resources of Washington, D.C., and Vicinity
GEOLOGICAL MAP
FIGURE 3-1

GEOLOGICAL MAP WASHINGTON PLATING WASHINGTON, D.C.

Scale 1:62,500



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Several other Quaternary age formations are exposed across the study area. The lowermost Quaternary age Sunderland Formation, consisting of orange-red to pink, yellow, and blue-gray coarse gravel, cross-bedded sand, silt, and clay, is exposed approximately 0.25 mile north of the site. The overlying Pamilico Formation, having a completely fluvial history, is exposed along the Potomac and Anacostia Rivers to the south. The Pamilico is composed of a mixture of gravel, sand, and silt-sized materials. Recent alluvial deposits of clay, sand, and gravel are present in the southwestern portion of the study area along the Potomac River. These alluvial deposits are generally only a few feet thick but may exceed 20 feet in some locales. 15

Scattered outcrops of the Tertiary age Bryn Mawr and Brandywine gravel deposits are present in the northern part of the study area. The Bryn Mawr gravel caps isolated hilltops and consists of coarse, poorly sorted pebbles in a red sand and silt matrix. The Brandywine gravel is pink or yellow and is composed of well-rounded, polished pebbles of quartzite, sandstone, and chert mixed with fairly clean quartz sand.¹⁵

The Cretaceous age Patapsco, Arundel, and Patuxent Formations, members of the Potomac Group, underlie the Tertiary sediments and overlie the Piedmont basement rocks. The upper Patapsco Formation consists of a basal maroon clay-rich layer that grades upward into a light-colored sandy zone. The total thickness of the Patapsco ranges between 200 and 500 feet. The Arundel Formation, underlying the Patapsco Formation, is a dark, tough clay that contains vast quantities of iron carbonate nodules and highly carbonaceous, lignitized tree trunks. The total thickness of the Arundel may exceed 200 feet. The Patuxent Formation, with a stratigraphic thickness between 140 and 300 feet, contains large percentages of sand that is commonly mixed with variable amounts of kaolin, mica, gravel, and lenses of variably colored or white, massive clay. 15,16

The oligoclase-mica facies of the lower Paleozoic age Wissahickon Formation is present in the northwestern part of the study area and is a garnet-rich quartz-muscovite schist of variable composition. The Wissahickon Formation has been intruded and altered by a number of igneous intrusives. The Kensington Granite Gneiss, Sykesville Formation, and Laurel Gneiss are granitic intrusives that have a schistose fabric, with many inclusions. The mafic igneous rocks that are encountered consist of tonalite, metadiorite, gabbro, amphibolite, and undifferentiated mafic rocks. The depth to this unit beneath the site is currently unknown.¹⁵

Because the entire site has been either paved or obstructed with buildings or other structures, the above information could not be confirmed during field operations. No lithologic exposures revealing bed orientation, presence of geologic contacts, or depth to bedrock were encountered.²

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3.3.2 Soils

Although soil maps for the site and study area are unavailable, it is considered, based upon the location of the site in a highly urbanized area, that the mapped materials are probably regarded as

Urban lands. Nearly all of the site is either covered with pavement or building structures. In several

unpaved areas adjacent to the site, a dark brown, sandy loam soil was encountered at the surface.

This material is probably a component of Urban land.²

3.3.3 Groundwater

Although essentially the entire population obtains potable water from public supplies derived from

surface water sources, a fairly good potential does exist for retrieval from groundwater origins.

In the crystalline rocks of the Piedmont Province, groundwater is stored and transmitted principally

along fractures, joints, cleavage planes, and bedding-plane separations and is controlled by the rock

structure. In the unconsolidated sediments of the Coastal Plain, groundwater movement is through

primary or intergranular porosity. 15

The Quaternary and Tertiary age sediments yield relatively small supplies of groundwater to shallow

wells. The Cretaceous age Patapsco and Patuxent Formations are generally the most productive

water-bearing formations in the Washington area. Yields obtained from the Patuxent range from 10

to 300 gallons per minute (gpm) and average 80 gpm. Wells tapping the Wissahickon Formation have

the greatest average yield for any of the Piedmont Formations, with yields ranging from 0.2 to 110

gpm. The wells supplying these yields are typically drilled to depths between 21 to 825 feet beneath

the surface. The average depth of wells across the study area is 124 feet, while the average yield is 13

gpm. Wells in formational contact have the highest yields, ranging from 5 to 40 gpm and averaging

16 gpm. 15

Generally, the water-table elevation reflects the local topography. Specific depths to the water table

are unknown for wells within the study area. Groundwater movement is downward and laterally

toward lower altitudes, eventually returning to the land surface through springs or wells. Recharge is

accomplished through precipitation and local bodies of surface water. Groundwater in the Piedmont

is typically considered to be unconfined or under water-table conditions, while groundwater in the

Coastal Plain sediments is encountered under both unconfined and confined, or artesian, conditions.

The direction of groundwater flow beneath the site is expected to be to the south, toward the

Potomac River. 15

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3.4 Climate and Meteorology

The average temperature for Laurel, Maryland which is 14.5 miles northeast of the site, is 55.9°F. The

average temperatures range from 33.4°F in January to 77.2°F in July. The average annual

precipitation is 41.96 inches. The mean annual lake evaporation is approximately 36.5 inches. The

net precipitation is approximately 5.46 inches. A 1-year, 24-hour rainfall will provided approximately

2.5 inches. 17,18,19

3.5 Land Use

The site lies within the District of Columbia, an urban area. Immediately surrounding the site, land

use is residential and commercial. The White House, the United States Capitol, and most of the

surrounding federal buildings lie one to three miles south of the site. Numerous public attractions,

such as the Smithsonian Institute, the Lincoln Memorial, and the Washington Monument also lie

within the three-mile radius of the site. The Henry Harrison Elementary School is immediately east of

the site. 1.2

3.6 <u>Population Distribution</u>

Approximately 125,000 people live within a 3-mile radius of the site. Approximately 56,340 people

live within a 2-mile radius. Approximately 12,520 people live within a 1-mile radius of the site. The

estimated number of persons living within a three-mile radius was obtained by estimating the area

percentage of the one-, two-, and three-mile radii within the District of Columbia. The population for

the District of Columbia was obtained from the 1980 Census Bureau data. 1,20

3.7 Critical Environments

One federally listed threatened crustacean is known to occur approximately 1.1 miles from the site in

Rock Creek. It is the Hay's Spring amphipoo (Stygobromus hayi). No critical habitats are located

within the site area.21

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4.0 WASTE TYPES AND QUANTITIES

The only wastes known to be on site are located in the flood water and sediment in the basement of

the old plating room. Analyses of samples taken from the basement on December 6, 1988 by NUS FIT

3 revealed the presence of heavy metals including lead (8,070 mg/kg), nickel (1,810 mg/kg), and zinc

(14,500 mg/kg), as well as semivolatiles and pesticides. All other tanks and drums of chemicals and

wastes that were used and generated on site have been moved to the new facility location at 2215

Adams Place Northeast, Washington, D.C.²

The first two tanks used in the electroplating process were tanks of hot caustic soda and acid used to

clean the bumpers prior to plating. The nickel-plating tanks used a solution of nickel sulfate, nickel

chloride, and boric acid. The two 1,500-gallon nickel-plating tanks were heated to 145°F. According

to the Baseline Report submitted to EPA, 12 drums of nickel sulfate and 2 drums of nickel chloride

were used, processed, or stored per year on the premises. The chromium-plating process used a 600-

gallon tank of chromic acid heated to 110°F. The chromic acid was purchased in 5-gallon fiber drums

with approximately 200 pounds on the premises at a time. 3,23

Rinse tanks were located after each of the nickel, chromium, and caustic soda tanks. Two of the 3

rinse tanks were 800 gallons and the other was 600 gallons. The tanks were drained into the public

sewer system two to three times each day through the center floor drain. The center floor drain also

collected any spills that occurred when bumpers were being moved from one tank to the next.

According to the Baseline Report, a total of 5 gpm or 2,400 gallons per day of rinse water were

drained into the public sewer system.3,22

Results from the initial samples obtained during the March 1986 overflow incident show high levels

of total chromium and total nickel. Total chromium levels of 70,600 ug/l and 78,300 ug/l were

detected in samples from the alley behind the plating building and in the alley behind Henry Harrison

School, respectively. Total nickel levels of 180,000 ug/l and 105,000 ug/l were detected, respectively,

in the same locations. These samples were aqueous effluent samples from puddles in the alleys after

the overflow incident occurred. 3,23

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5.0 FIELD TRIP REPORT

5.1 Summary

On Thursday, December 6, 1988, FIT 3 members Elizabeth Coughlin, Thomas Bachovchin, Robert Chappell, Scott Coslett, and Michael McCarthy visited the Washington Plating site in Washington, D.C. FIT 3 met with George Galich and Jose Flores, of Washington Plating. Weather conditions during the site visit were sunny, with temperatures in the mid-50s.

One aqueous and one sediment sample were obtained on site (see figure 5.1, page 5-4). Five off-site soil samples were obtained. No split samples were requested. Photographs were taken on site (see figure 5.2, page 5-6, and the photograph log, section 5.5).

Deviations from the Sampling Plan

There were no deviations from the sampling plan.

5.2 Persons Contacted

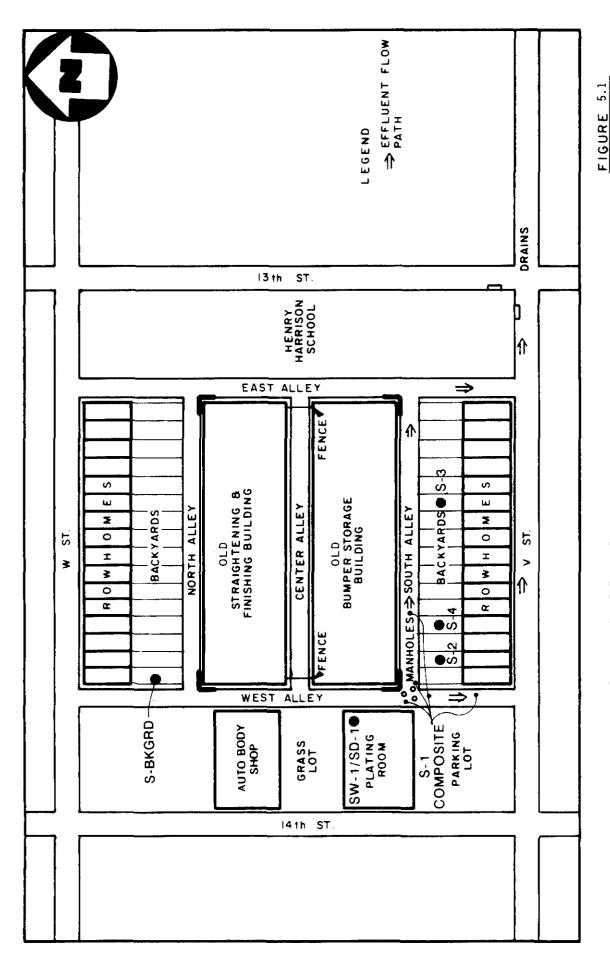
5.2.1 Prior to Field Trip

George Galich President Washington Plating Company 2215 Adams Place Northeast Washington, DC 10018 (202) 636-8715

Byron Bacon
Department of Consumer and Regulatory Affairs
613 G Street Northwest
Washington, DC 20004
(202) 783-3193

Milan Milosevic Manager Washington Plating Company 2215 Adams Place Northeast Washington, DC 10018 (202) 636-8715

James McCreary
Site Investigation Officer
U.S. EPA
841 Chestnut Building
Ninth and Chestnut Street
Philadelphia, PA 19107
(215) 597-1105



WASHINGTON PLATING CO., WASHINGTON, D.C.

A Halliburton Company

CORPORATION

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5.2.2 At the Site

George Galich President 2215 Adams Place Northeast Washington, DC 10018 (202) 636-8715

5.2.3 Post Site Visit

Milan Milosevic Manager Washington Plating Company 2215 Adams Place Northeast Washington, DC 10018 (202) 636-8715 Jose Flores Employee Washington Plating Company 2215 Adams Place Northeast Washington, DC 10018 (202) 636-8715

James McCreary
Site Investigation Officer
U.S. EPA
841 Chestnut Building
Ninth and Chestnut Streets
Philadelphia, PA 19107
(215) 597-1105

5.2.4 Water Supply Well Information

There are no know home wells within three miles of the site.

TDD NUMBER <u>F3-8810-15</u>

EPA NUMBER <u>DC · CC7</u>

5.3 SAMPLE LOG

SITE NAME WASHINGTON PLATING

	TRAFFIC REPORTS		SAMPLE IDENTIFIER	PHASE	SAMPLE DESCRIPTION	SAMPLE LOCATION	TARGET USE	pН	FIELD MEASUREMENTS
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C Y	MCY	4320-0 2	5-2	مور،	Pt FIR Supples see		Accessable to me Albuc.		,
CY C71	MC7 C70	4320-C- 3	5.3	Soc.	Off SINE SURFACE SULL DAME AICH BREWN LLAMY SULL		ACLESSABLE TO THE RESIDENTS DO 1331 V SMETT		
CY C72	MC1 	4326-C=	5-4	Søc.	Off SITE SURFACE SOIL		ACCESSABLE TO THE MESTORNES OF 1343 V STANCET		İ
CY 073	MCY C-72		5-5a	Soc	Dupuna 25-1		Deputerte of 5.1		
C Y c 74	MC7 073	4320-C- L	S-BACK	کور	BACKGARUND BACK BREWN LOANY SHE WITH JAMPS TO		Accessable TO THE PUBLIC		HALL BACK EROUND READING WAS D.2 PPM, THE MAGATION MINE ALBER WAS SET AT X1; WE REMINES ABOVE DACKGEMIND.
(4	MC 7	4325 C	Sa (5	ON SITE SUDEMENT SAMOY BLACK SITE WITH CALBACK.		ACCUTES RESTRICTED		
(()	() () ()	11/4	jo tar e t	Ai	CN SIR SURFACE UNDER CLORE		ALLESS LESTRICTED	6.33	
27	Mc/	N/A	Ay PLANK	Aa	ELME	BLANE	BLANK		2

TDD NUMBER F3-8810-15

EPA NUMBER DC - 007

5.3 SAMPLE LOG

SITE NAME WASHINGTON PLATING

TR Organi c	AFFIC REPO	ORTS High Hazard	SAMPLE IDENTIFIER	PHASE	SAMPLE DESCRIPTION	SAMPLE LOCATION	TARGET USE	ρН	FIELD MEASUREMENTS
CY C78	N/A	N/A	S BLANK	Aq	BLANK	BLANK	BLANK		
									,
	-								
									E.
									3

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5.4 <u>Site Observations</u>

- The HNU background reading was 0.1 ppm. No readings above background were recorded.
- The radiation mini-alert was set at the X1 position. No readings above background were recorded.
- The water in the flooded basement of the plating building was approximately four feet deep. The basement was approximately 65 feet long and 10 feet wide.
- No stains were observed in either the alley or the soils in neighboring backyards.
- All buildings were locked at the time of the FIT visit.
- The center alley was fenced.
- No drums were found on site.

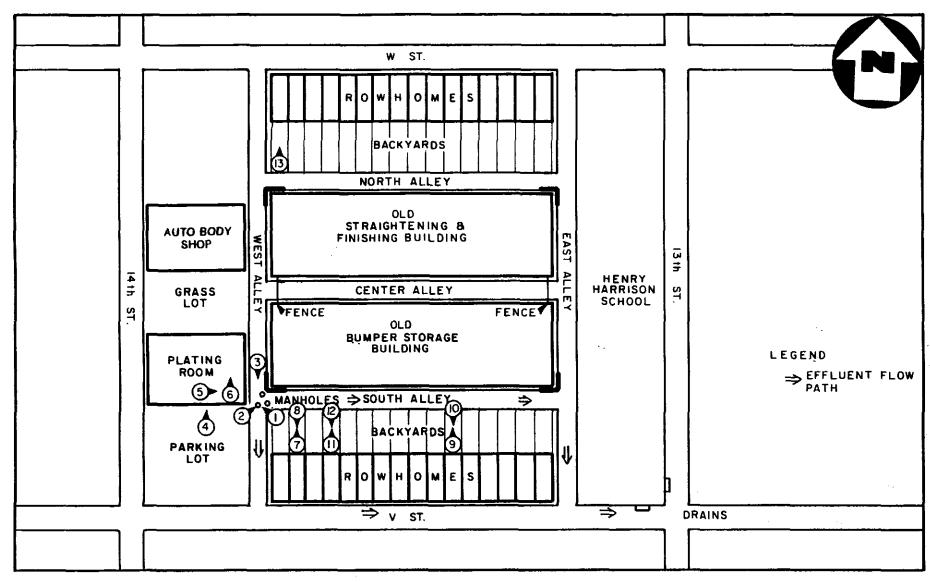


PHOTO LOCATION MAP

WASHINGTON PLATING CO., WASHINGTON, D.C.

(NO SCALE)

FIGURE 5.2



EPA REGION III SUPERFUND DOCUMENT MANAGEMENT SYSTEM

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EPA

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 1 - SITE LOCATION AND INSPECTION INFORMATION

I. IDENTIFICATION

01 STATE 02 SITE NUMBER

PA	RT 1 - SITE LO	CATION AND IN	SPECTION INFO	RMATION UTS	007
II. SITE NAME AND LOCATION					
or SITE NAME (Legal, common, or descriptive na Washington Plating	ume of site)		02 STREET, ROUTE NO., or 2109 14th Stree	n specific Location identifier et Northwest	•
oscity Washington			04 STATE 05 ZIP CODE 20009	06 COUNTY N/A	07 COUNTY CODE 08 CONG. DIST DC01
09 COORDINATES LATITUDE 38° 35¹ 06"	LONGITUDE 77° 01' 55"	10 TYPE OF OWNER A. PRIVATE F. OTHER		C. STATE	O COUNTY E. MUNICIPAL
III. INSPECTION INFORMATION					
01 DATE OF INSPECTION 12/6/88	02 SITE STATUS ACTIVE INACTIVE		973 L	1987	UNKNOWN
04 AGENCY PERFORMING INSPECTION (Check all	that apply) NUS Corporati (Name of firm) O		HCIPAL CONTRACTOR (Specify)	(Name of firm)
05 CHIEF INSPECTOR	06 TIT	rue ologist		07 ORGANIZATION NUS FIT 3	08 TELEPHONE NO. (215) 687-9510
09 OTHER INSPECTORS (b) (4)	10 717			11 ORGANIZATION NUS FIT 3	12 TELEPHONE NO. (215) 687-9510
(b) (4)	Ge	ologist		NUS FIT 3	(215) 687-9510
(b) (b) (4)	Ge	ologist		NUS FIT 3	(215) 687-9510
(b) (4)	En	vironmental Tec	chnician	NUS FIT 3	(215) 687-9510
13 SITE REPRESENTATIVES INTERVIEWED George Galich	14 ftf	esident	is accass, Washing 2215 Adams Plac Washington, DO		16 TELEPHONE NO (202) 636-8715
Jose Flores	Em	ployee	Washington Plan 2215 Adams Plan Washington, DC	ting Company	(202) 636-8715
(Charles and	inspection 00 a.m.	19 WEATHER CONDITION Sunny, with	temperatures in	the 50s.	
IV. INFORMATION AVAILABLE FROM	l				
OICONTACT James McCreary		(Agency/Organization) S. EPA			03 TELEPHONE NO (215) 597-1105
04 PERSON RESPONSIBLE FOR SITE INSPECTION ((b) (4)	FORM 05 AC	iency IS	06 ORGANIZATION FIT 3	07 TELEPHONE NO. (215) 687-9510	08 DATE 6/30/89



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 2 - WASTE INFORMATION

i	I. IDENTIFIC	ATION
	01 STATE	02 SITE NUMBER
-	00	007

II. WASTE STA	TES, QUANTITIES, AND CH	ARACTERIS	TICS					PROPERTY.
A SOLID B POWD C SLUDG D OTHER	ER, FINES F. LIQUID E G. GAS	(Mess indep			O3 WASTE CHAI A. TOXIC B. CORROSIVE C. RADIOACTIVE D PERSISTENT		E SOLUGLE HINFECTIOUS FLAMMABLE HIGHITABLE	THE ADDITION OF THE
III. WASTE TŸF	PE							
CATEGORY	SUBSTANCE NAM	E	01 GROSS AMOUNT	02 UNI	T OF MEASURE	03 CC	OMMENTS	
SLU	SLUDGE							
OLW	OILY WASTE			†		<u>† </u>	000	
SOL	SOLVENTS							
PSD	PESTICIDES				-	1		
occ	OTHER ORGANIC CH	EMICALS						
ЮС	INORGANIC CHEMIC	ALS		†	· · · · · · · · · · · · · · · · · · ·	†		
ACD	ACIDS					t	-	
8AS	BASES			† 	• 0	1		
MES	HEAVY METALS					<u> </u>		
/. HAZARDOL	JS SUBSTANCES (See Appendi	t for most free	quently cited CAS Numbers)	_				
OT CATEGORY MES	- 02 SUBSTANCE NAME		03 CAS NUMBER 7439-97-6	04 STO Spill:	PRAGE DISPOSAL MET	ноо	05 CONCENTRAT	ION 06 MEASURE OF CONCENTRATION
MES	chromium		7740-47-3	Spills			2,900	mg/kg
MES	lead		7439-92-1	Spill			8,070	mg/kg
MES	zinc		7440-66-6	Spill			14,500	mg/kg
MES	cyanide		57-12-5	Spill:			10.60	mg/kg
OCC	naphthalene		91-20-3	Spill:			1,700	ug/kg
000	acenaphthylene		208-96-8	Spill:		-	370	ug/kg
occ	dibenzofuran		132-64-9	Spill:			460	ug/kg
000	pentachlorophenol		87-86-5	Spill:			5,600	ug/kg
0CC	bis(2-ethylhexyl) ph	thalate	117-81-7	Spill:			23,000	ug/kg
								ug/kg
V. FEEDSTOCI	KS (See Appendix for CAS Numbers 01 FEEDSTOCK NA		02 CAS NUMBER	CATEG	ORY 016	EEDST	OCK NAME	02 CAS NUMBI
FDS				FDS				
FDS				FDS	<u> </u>			
				FDS				
FDS			•					_1
				FDS	3			Ī



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 3 - DESCRIPTION OF HAZARDOUS CONDITIONS AND INCIDENTS

1	I. IDENTIFIC	ATION
	O1 STATE	02 SITE NUMBER

II. HAZARDOUS CONDITIONS AND INCIDENTS				
01 A. GROUNDWATER CONTAMINATION 03 POPULATION POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE:) [POTENTIAL	☐ ALLEGED
01 8. SURFACE WATER CONTAMINATION 03 POPULATION POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE: 04 NARRATIVE DESCRIPTION) 🗖	POTENTIAL	ALLEGED
01 C. CONTAMINATION OF AIR 03 POPULATION POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE: 04 NARRATIVE DESCRIPTION) 🗖	POTENTIAL	☐ ALLEGED
01 D FIRE/EXPLOSIVE CONDITIONS 03 POPULATION POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE: 04 NARRATIVE DESCRIPTION) [POTENTIAL	ALLEGED
01 E. DIRECT CONTACT 03 POPULATION POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE:04 NARRATIVE DESCRIPTION) [POTENTIAL	ALLEGED
01 F CONTAMINATION OF SOIL 03 AREA POTENTIALLY AFFECTED: <0.25 (Acres) A soil sediment sample from the basement is and elevated levels of inorganic contaminations.		-		☐ ALLEGED taminants
01 G. DRINKING WATER CONTAMINATION 03 POPULATION POTENTIALLY AFFECTED: None reporte or observed	02 GOBSERVED (DATE:04 NARRATIVE DESCRIPTION) 0	POTENTIAL	☐ ALLEGED
01 H. WORKER EXPOSURE/INJURY 03 WORKERS POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE:) 🗖	POTENTIAL	☐ ALLEGED
01 POPULATION EXPOSURE/INJURY 03 POPULATION POTENTIALLY AFFECTED: None reported or observed	02 OBSERVED (DATE:) [POTENTIAL	ALLEGED



POTENTIAL HAZARDOUS WASTE SITE SITE SITE INSPECTION REPORT PART 3 - DESCRIPTION OF HAZARDOUS CONDITIONS AND INCIDENTS

I. IDENTIFIE	CATION
01 STATE	ON WEST PROPERTY.

N. HAZARDOUS CONDITIONS AND INCIDENTS (Continue	od)	
01 DAMAGE TO FLORA 04 NARRATIVE DESCRIPTION	02 GOBSERVED (DATE:)	O POTENTIAL O ALLEGED
None reported or observed		
01 K DAMAGE TO FAUNA 04 NARRATIVE DESCRIPTION (Include name(s) of specie None reported or observed	02 GOBSERVED (DATE:) es)	D POTENTIAL D ALLEGED
01 □ L. CONTAMINATION OF FOOD CHAIN 04 NARRATIVE DESCRIPTION None reported or observed	02 GOBSERVED (DATE)	☐ POTENTIAL ☐ ALLEGED
01 M UNSTABLE CONTAINMENT OF WASTES (Spills, Runoff, Standing liquids, Leaking drug 12,520 wi 03 POPULATION POTENTIALLY AFFECTED: 1 mile	02 GOBSERVED (DATE: 12/6/88) thin 04 NARRATIVE DESCRIPTION	POTENTIAL ALLEGED
There was an observed chemical spill in to volatile organic contaminants and elevate	 the basement. Sample results indicate	ed elevated levels of semi-
01 O N DAMAGE TO OFF-SITE PROPERTY 04 NARRATIVE DESCRIPTION	02 GOBSERVED (DATE:)	POTENTIAL ALLEGED
None reported or observed		
01 G O CONTAMINATION OF SEWERS, STORM ORAINS, WWTPS 04 NARRATIVE DESCRIPTION None reported or observed	02 GOBSERVED (DATE:)	☐ POTENTIAL ☐ ALLEGED
01 P ILLEGALUNAUTHORIZED DUMPING 04 NARRATIVE DESCRIPTION None reported or observed	02 GOSERVED (DATE:)	POTENTIAL ALLEGED
05 DESCRIPTION OF ANY OTHER KNOWN, POTENTIAL	, OR ALLEGED HAZARDS	
None reported or observed		
II. TOTAL POPULATION POTENTIALLY AFFECTED:12.	520 within 1 mile	
V. COMMENTS		
Rainfall has caused an overflow from the	basement of the former facility onto	private property.
V. SOURCES OF INFORMATION (Cite specific references,	e.g., state files, sample analysis, reports)	
NUS FIT 3. Site inspection. TDD No. F3	3-8810-15, December 6, 1988.	

E	P	A
	E	EP

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 4 - PERMIT AND DESCRIPTIVE INFORMATION

1	Ł.	DENTIFK	ATI	OM	
ſ	01	STATE	02	SITE	NUMBER

(Red)

					THE STATE OF THE S		
H. PERMIT INFORMATION					(Red)		
01 TYPE OF PERMIT ISSUED (Check all that apply)			03 DATE ISSUED 04 EXPIRATION DATE		05 COMMENTS		
A. NPDES	1						
☐ B UIC							
C. AIR		Î					
D RCRA							
E. RCRA INTERIM STATUS							
F SPCC PLAN							
☐ G: STATE (Specify)							
☑ H LOCAL(Specify) District of Columbia	N/A	12/86	1/87	se	ewer discharge		
☐ I. OTHER (Specify)							
☐ J. NONE							
III. SITE DESCRIPTION							
G1 STORAGE DISPOSAL (Check all that apply)	02 AMOUNT 03 UNIT OF M	MEASURE 04 TREAT	MENT (Check all that apply)	05 OTHER		
☐ A. SURFACE IMPOUNDMENT ☐ B. PILES ☐ C. DRUMS, ABOVE GROUND ☐ D. TANK, ABOVE GROUND ☐ E. TANK, BELOW GROUND		☐ B. UNDERGROUND INJECTION ☐ C. CHEMICAL/PHYSICAL ☐ D. BIOLOGICAL ☐ E. WASTE OIL PROCESSING			D A. BUILDINGS ON SITE		
F. LANDFILL G. LANDFARM H. OPEN DUMP SI OTHER	□ F. \$O\ □ G. OT	☐ F. SOLVENT RECOVERY ☐ G. OTHER RECYCLING/RECOVERY ☐ N/A (Specify) □ DH. OTHER N/A (Specify)					
07 COMMENTS Around 1986, a blockage in a sewer line caused an overflow of plating baths and/or rinses to be discharged into the basement of the plating room. Rain water has added to the volume of the spill.							
IV. CONTAINMENT							
01 CONTAINMENT OF WASTES (Check of A. ADEQUATE, SECURE		C. INADEQUATE,	POOR DI	NSECURE,	UNSOUND, DANGEROUS		
02 DESCRIPTION OF DRUMS, DIKING, LINERS, BARRIERS, ETC. The spilled plating baths and/or rinses are located in a concrete basement.							
V. ACCESSIBILITY							
01 WASTE EASILY ACCESSIBLE: YE 02 COMMENTS Waste is located in the basem	_	lding.					
VI. SOURCES OF INFORMATION (Cite specific references, e.g., state files, sample analysis, reports)							
NUS FIT 3. Site inspection.	TDD No. F3-8810-15,	, December 6, 1	988.				

EPA

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 5 - WATER, DEMOGRAPHIC, AND ENVIRONMENTAL DATA

I. IDENTIFICATION					
01 STATE	02 SITE NUMBER				

PART	5 - WATER,	DEMOGRAPHI	C, AND ENVIR	ONMENTA	LDATA	DC DC	9846	NUMBER ///A	
II. ORINKING WATER SUPPLY									
01 TYPE OF DRINKING SUPPLY		02 STATUS				0	3 DISTANCE TO	SITE	
(Check as applicable) SURFACE	WELL	ENDANGERED	AFFECTED	MONITO	DRED] , .	13.6	(m.)	
COMMUNITY A. M	B. 🗖	6 :□	8 - E -	С. <u>.</u> Б		<u> </u>	13.0	(mi)	
					<u> </u>				
III. GROUNDWATER 01 GROUNDWATER USE IN VICINI	TV /Ch - ch - c- c						·		
☐ A ONLY SOURCE FOR ORINKING	☐ 8. ORIN (Othi COM		(Li IRIGATION	AMERCIAL, INDU mited other sour		rion (NOT USED.	UNUSABLE	
02 POPULATION SERVED BY GROUNDWATE	a <u>0</u>		03 DISTANCE TO	O NEAREST DRIN	KING WATER V	rell N/A		(mi)	
04 DEPTH TO GROUNDWATER	05 DIRECTION	OF GROUNDWATER FLOY	OS DEPTH TO A		07 POTENTIA OF AQUI		00 SOLE SOU	RCE AQUIFER	
<u> <124</u> (ft)	to th	e south	: i		115,200		☐ YES	₽ NO	
124 feet. Specific well 10 RECHARGE AREA COMMENTS			11 DISCHARGI	E AREA	; Groundw	ater prob	ably disc	harges	
Recharge is acco	-	•	tion 🗖 NO		into th	e Potomac	River.		
IV. SURFACE WATER		<u>-</u>							
01 SURFACE WATER USE (Check o	□ B. HRRK	GATION, ECONOMICALLY PORTANT RESOURCES	□c. con	IMERCIAL, INDU	STRIAL	ſ	D. NOT CURRE	ENTLY USED	
02 AFFECTED/POTENTIALLY AFFE	CTED BODIES	OF WATER							
NAME:				AFFECTED			DISTANCE TO SITE		
N/A		- 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1						(mı)	
					_			(mi) (mi)	
V. DEMOGRAPHIC AND PROPERTY	INSCREASE.	W							
01 TOTAL POPULATION WITHIN				1				· · · · ·	
	(2) MILES OF	SITE THREE!	B) Miles OF SITE	02 015	STANCE TO NEA	REST POPULATI	ON		
A 12,520 B	56,340	,	25,200 NO. OF PERSONS		200	feet	_ (111 1)		
D3 NUMBER OF BUILDINGS WITHIN TWO (2)	MILES OF SITE	040	ISTANCE TO NEAREST O	FF-SITE BUILDING	<u> </u>				
14,826				20	feet	_(mı)			
05 POPULATION WITHIN VICINITY OF SITE (P The site is located in Wa	rovide narrative de	Reciption of nature of po	pulation within vicinity	of site, e.g., rural	i, village, dense	ly populated urt	ban area)		



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 5 - WATER, DEMOGRAPHIC, AND ENVIRONMENTAL DATA

I. IDENTIFICATION (Red)

VI. ENVIRONMENTAL INFORMATIO				III DEE		
01 PERMEABILITY OF UNSATURATE						
_	•	C. 10 ⁻⁴ - 10 ⁻³ cm/sec	D. GREATER T	'HAN 10 ⁻³ cm/sec		
02 PERMEABILITY OF BEDROCK (Che A. IMPERMEABLE 18. (Less than 10-5 cm/sec)	eck one) $10^{-3} - 10^{-5}$ cm RELATIVELY IMPERMEABLE C (10-1-10-1 cm/sec)	/sec IC. RELATIVELY PEF (10-2 - 10-4 cm/se	RMEABLE D. VERY F C) (Great	PERMEABLE ter than 10-2 cm/sec)		
03 DEPTH TO BEDROCK	04 DEPTH OF CONTAMINATED SOIL ZONE	05 SOIL pH	known			
<u><124</u> (ft)	unknown(ft)		KIIOWII			
DE NET PRECIPITATION	07 ONE-YEAR 24-HOUR RAINFALL 2.5 (in)	OB SLOPE SITE SLOPE	DIRECTION OF SITE SLOPE	LOPE TERRAIN AVERAGE SLOPE		
(in)	(IR)	_<1%	southeast			
SITE IS IN N/A YEAR FLOOD		ARRIER ISLAND, COASTAL	. HIGH HAZARD AREA, RIVERINE FL	OGDWAY		
11 DISTANCE TO WETLANDS (S-acre minimum) ESTUARINE A 3/4 (mi)	OTHER(mi)		CRITICAL HABITAT (of endangered N/A EREO SPECIES:			
DISTANCE TO: COMMERCIAL/INDUSTRIA A 50 feet (500)		LIFE RESERVES	PRIME AG LAND C. >3. (mi)			
14 DESCRIPTION OF SITE IN RELATION The site lies on the broad	ON TO SURROUNDING TOPOGRAP		er, 2.5 miles north o	of the river.		
VII. SOURCES OF INFORMATION (Gre						

Johnston, P.M., United States Geological Survey. Geology and Groundwater Resources of Washington, D.C. and

Vicinity. Geological Survey Water Supply Paper 1776, undated.



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 6 - SAMPLE AND FIELD INFORMATION

i. IDENTIFIC	ATION
01 STATE	02 SITE NUMBER

SAMPLE TYPE SINDAMES TAKEN GROUNDWATER Organic samples were sent to IT SURFACE WATER 1 WASTE Inorganic samples were sent to JTC AIR RUNOFF Hexavalent chrome samples were sent to Chen Tech SPILL SOIL 7 VEGETATION OTHER Blank 2 IM. FIELD MEASUREMENTS TAKEN OI TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were meteroaded. Where meteroaded. Radiation mini-alert DATE W. PHOTOGRAPHS AND MAPS OI TYPE O3 GROUND AERIAL O2 IN CUSTODY OF NUS Corporation (Name of organizacor or meteroqual) O3 MAPS ON VOTHER FIELD DATA COLLECTED (Provide marries we described) N/A VI. SOURCES OF INFORMATION (cits seedlic reference, i.g., Nate fifes, sample analysis, reports)	ed;
SURFACE WATER I Inorganic samples were sent to II AIR RUNOFF Hexavalent chrome samples were sent to Chen Tech SPILL SOIL 7 VEGETATION OTHER Blank 2 IF FIELD MEASUREMENTS TAKEN IT TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The minitalent was set at the XI position. No readings above background were recorded. V. PHOTOGRAPHS AND MAPS O1 TYPE O3 GROUND AERIAL O2 IN CUSTODY OF NUS Corporation (Name of organization or neurolus) NAA OTHER FIELD DATA COLLECTED (Promise neurolus described) N/A	DATE /AILABLE
SURFACE WATER WASTE Inorganic samples were sent to JTC AIR RUNOFF SPILL SOIL 7 VEGETATION OTHER Blank 2 INTELD MEASUREMENTS TAKEN 1TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. HNU The mini-alert was set at the X1 position. No readings above background were recorded. IntelD MEASUREMENTS TAKEN 1 TYPE O3 GROUNO AERIAL O2 IN CUSTODY OF NUS Corporation Thanked organisation or individual The mini-alert was set at the X1 position. No readings above background were recorded. O HOTOGRAPHS AND MAPS TYPE O3 GROUNO AERIAL O2 IN CUSTODY OF NUS Corporation Thanked organisation or individual THE STATE OF TH	<u> </u>
WASTE Inorganic samples were sent to JTC AIR RUNOFF Hexavalent chrome samples were sent to Chem Tech SPILL SOIL 7 VEGETATION OTHER Blank 2 INIELD MEASUREMENTS TAKEN ITYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. HNU Radiation mini-alert MAPS O4 LOCATION OF MAPS O3 YES NUS FIT 3 Site inspection. TDD No. F3-8810-15 OTHER FIELD DATA COLLECTED (Provide Nurstave description) N/A	
AIR RUNOFF SPILL SOIL 7 VEGETATION OTHER Blank 2 I. FIELD MEASUREMENTS TAKEN 1 TYPE 02 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. I. PHOTOGRAPHS AND MAPS 1 TYPE 12 GROUND AERIAL 13 MAPS 14 VES NO NO NO NUS FIT 3 Site inspection. TDD No. F3-8810-15. COTHER FIELD DATA COLLECTED (Provide Narrative description) N/A	
SPILL SOIL 7 VEGETATION OTHER Blank 2 I. FIELD MEASUREMENTS TAKEN 1 TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. V. PHOTOGRAPHS AND MAPS 1 TYPE O2 GROUND AERIAL O2 IN CUSTODY OF NUS Corporation (Name of organization or individual) WAPS O3 YES NUS FIT 3 Site inspection. TDD No. F3-8810-15	<u></u>
VEGETATION OTHER Blank 2 I. FIELD MEASUREMENTS TAKEN I TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. I. PHOTOGRAPHS AND MAPS I TYPE O3 GROUND AERIAL O2 IN CUSTODY OF NUS Corporation (Name of organization or individual) O3 MAPS O4 LOCATION OF MAPS O3 YES NUS FIT 3 Site inspection. TDD No. F3-8810-15 O OTHER FIELD DATA COLLECTED (Provide narrative description)	
VEGETATION OTHER Blank 2 I. FIELD MEASUREMENTS TAKEN I TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded. If the mini-alert was set at the X1 position. No readings above background were recorded.	
OTHER Blank 2 I. FIELD MEASUREMENTS TAKEN ITYPE 02 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. I. PHOTOGRAPHS AND MAPS ITYPE 03 GROUND AERIAL 02 IN CUSTODY OF NUS Corporation (Name of organization or individual) 3 MAPS 04 LOCATION OF MAPS O2 YES NO NUS FIT 3 Site inspection. TDD No. F3-8810-15	
I TYPE O2 COMMENTS The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. A. PHOTOGRAPHS AND MAPS 1 TYPE O2 GROUND AERIAL O2 IN CUSTODY OF NUS Corporation (Name of organization or individual) 3 MAPS O4 LOCATION OF MAPS O2 YES NUS FIT 3 Site inspection. TDD No. F3-8810-15 OTHER FIELD DATA COLLECTED (Provide narrative description)	
The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. 7. PHOTOGRAPHS AND MAPS 1 TYPE	
The background reading was 0.1 ppm. No readings above background were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. PHOTOGRAPHS AND MAPS TYPE GROUND AERIAL 02 IN CUSTODY OF NUS Corporation (Name of organization or individual) 3 MAPS AUGUSTION OF MAPS GYES NUS FIT 3 Site inspection. TDD No. F3-8810-15 OTHER FIELD DATA COLLECTED (Frounds narrative description)	
HNU were recorded. The mini-alert was set at the X1 position. No readings above background were recorded. F. PHOTOGRAPHS AND MAPS 1 TYPE	
TYPE S GROUND AERIAL 02 IN CUSTODY OF NUS Comporation (Name of organization or individual) 3 MAPS S O4 LOCATION OF MAPS NUS FIT 3 Site inspection. TDD No. F3-8810-15 OTHER FIELD DATA COLLECTED (Provide narrative description)	ru
1 TYPE	
1 TYPE GROUND AERIAL 02 IN CUSTODY OF NUS Corporation (Name of organization or individual) 3 MAPS	
AMAPS □ YES □ NO NUS FIT 3 Site inspection. TDD No. F3-8810-15 OTHER FIELD DATA COLLECTED (Provide narrative description) N/A	
3 MAPS ☐ YES ☐ NO	
NO NUS FIT 3 Site inspection. TDD No. F3-8810-15 OTHER FIELD DATA COLLECTED (Provide narrative description) N/A	
N/A	
N/A	
/I. SOURCES OF INFORMATION (Cite specific references, e.g., state files, sample analysis, reports)	
/L SOURCES OF INFORMATION (Cite specific references, e.g., state files, sample analysis, reports)	
/I. SOURCES OF INFORMATION (Cite specific references, e.g., state files, sample analysis, reports)	
/I. SOURCES OF INFORMATION (Cite specific references, e.g., state files, sample analysis, reports)	
NUS FIT 3. Site inspection. TDD No. F3-8810-15, December 6, 1988.	
100 (1) 0: 0100 (Hapecolon, 100 No. 15-0010-13, December 0, 1900.	



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT

I.	DENTIFIC	ΆΤΚ) (1	
		_		
D:	STATE	כמ ל	UTE	

			PART 7 - OW	NER INFORMATION		<u>. </u>		007
H. CURRENT OWNER(S)				PARENT COMPANY (if applicable)				
George Galich		+ 8 NUMBER	OS NAME N/A			09 D	+ 8 NUMBER	
03 STREET ADDRESS (P.O. Sox, NFD P. etc.) 2215 Adams Place Northeast			04 SIC CODE	10 STREET ADDRESS (P.O. Box, RFD #, etc.)	10 STREET ADDRESS (P.O. Box, RFD #, etc.)			11 SIC CODE
05 CITY Washington	OG STATI		17 ZIP CODE 10018	12 CITY		13 STATE	Ţ	4 ZIP CODE
OI NAME	<u> </u>	02 0	+ 8 NUMBER	08 NAME N/A			09 D - 8 NUMBER	
03 STREET ADDRESS (P.O. Box, RFD €, etc.)			04 SIC COD€	10 STREET ADDRESS (P.O. Box, RPD #, etc.)				11 SIC CODE
OS CITY	06 STATE	. (17 ZIP CODE	12 CITY		13 STATE	1	4 ZIP CODE
01 NAME		02 0	+ 8 NUMBER	QB NAME N/A			090	+ 8 NUMBER
03 STREET ADDRESS (P.O. Box. RFD #, etc.)			04 SIC COD€	10 STREET ADDRESS (P O. Box, RFD #, etc.)		- "		11 SIC COD€
05 CITY	06 STATE		7 ZIP CODE	12 CITY		13 STATE	T	4 ZIP CODE
01 NAME	II NAME 021		+ 8 NUMBER	00 NAME N/A	GB NAME N/A		09 D + 8 NUMBER	
03 STREET ADDRESS (P.O. Box, RFD #, etc.)			04 SIC CODE	10 STREET ADDRESS (P.O. BOX, RFD #, etc.)				11 SC CODE
DS CITY 06 STATE 07		7 ZIP CODE	12 CITY		13 STATE	1	4 ZIP CODE	
III. PREVIOUS OWNER(S) (list most recent	first)			IV. REALTY OWNER(5) (if applicable,	list mo	st recen	t fir	st)
01 NAME 02 D + 8 NUMBER			+ 8 NUMBER	01 NAME N/A		1	02 D	- 8 NUMBER
03 STREET ADDRESS (P.O. Box, RFD ♣, etc.)			04 SIC CODE	03 STREET ADDRESS (P.O. Box, AFD #, etc.)				04 SIC COD€
05 CITY	06 STATE		7 ZIP CODE	05 CITY		OG STATE	lo	7 ZIP CODE
01 NAME		02 D	+ 6 NUMBER	01 NAME N/A		()2 O	+ 8 NUMBER
03 STREET ADDRESS (P. O. Box, RFD. #, etc.)			04 SIC CODE	03 STREET ADDRESS (P.O. Box, RFD #, etc.)		L	3	04 SIC CODE
05 CITY	06 STATE	T	7 ZIP CODE	05 CITY		OS STATE	To	7 ZIP CODE
01 NAME		02 D	+ 8 NUMBER	01 NAME N/A		(12 0	- a NUMBER
03 STREET ADDRESS (P.O. Box, 1890 4, etc.)			04 SIC CODE	03 STREET ADDRESS (P.O. Box, AFD #, etc.)			04 SIC CODE	
05 CITY	06 STATE	. 0	7 ZIP CODE	05 CITY		06 STATE	0	ZIP CODE
V. SOURCES OF INFORMATION (Cite specific	eferences,	• g s	tate files, sample and	Hysis, reports)				
NUS FIT 3. Site inspection.			<u> </u>					
noo iii oi oi oo iiispeeeioii.	100 11	0. 1	5-0010-15 , L	ecember 0, 1900.				

EPA

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 8 - OPERATOR INFORMATION

I. IDENTIFICATION							
01 STATE	02	SITE TOWNSDER					
DC	L	/LQ07					

								1100
H. CURRENT OPERATOR (Pr	ovide if different	from a	wne:	r)	OPERATOR'S PARENT COMPANY (if a	oplicable)		
OINAME Washington Plating Company			02 0 +	8 NUMBER	10 NAME N/A		11 0 + 8 NUMBER	
03 STREET ADDRESS (P.O. Cox. NºO 2215 Adams Place I				04 SIC CODE	12 STREET ADDRESS (P.O. Box, RFD #, etc.)			13 SIC CODE
oscity Washington		DC		zi p coo€ 10018	14 CITY	15 STATE	,	6 ZIP COOE
08 YEARS OF OFERATION 09 NAME OF OWNER 10 George Galich								
III, PREVIOUS OPERATOR(S) (List mo	ost recent first; provide	only if di	fferen	t from owner)	PREVIOUS OPERATORS' PARENT COM	PANIES (if ag	plic	able)
01 NAME Unknown			02 D +	6 NUMBER	10 NAME N/A		110	+ B NUMBER
03 STREET ADDRESS (F.O. Box, RFD	ADDRESS (P.O. Box, RFD. Ø, etc.) 04 SIC CODE 12 STREET ADDRESS (P.O. Box, RFD. Ø, etc.)		12 STREET ADDRESS (P.O. Box, RFD #, etc.)) #, etc.)		13 SIC CODE		
05 CITY	0	6 STATE	07	ZIP CODE	14 CITY	15 STATE	,	6 ZIP CODE
08 YEARS OF OPERATION	09 NAME OF OWNER	DURING	THIS PE	ERIOD				
01 NAME 02 D + 8 NUMBER			8 NUMBER	10 NAME N/A			+ B NUMBER	
03 STREET ADDRESS (P.O. Box, RFD	ø. etc.)			04 SIC CODE	12 STREET ADDRESS (P.O. Box, RFD #, etc.)			13 SIC CODE
35 CITY	o	6 STATE	07	ZIP CODE	14 CITY	15 STATE	T	6 ZIP CODE
08 YEARS OF OPERATION	09 NAME OF OWNER	DURING	THS PE	RIOD				<u>· ·</u>
01 NAME 02 D + B N			B NUMBER	10 NAME N/A 11 D			+ 8 NUMBER	
D3 STREET ADDRESS (P.O. Box, RFD #, etc.) Q4 SIC COD€			12 STREET ADDRESS (P.O. Box, RFD #, etc.)			13 SIC CODE		
05 CITY	0	6 STATE	07	ZIP CODE	14 CITY	15 STATE	T	6 ZIP CODE
OB YEARS OF OPERATION OF NAME OF OWNER DURING THIS PERIOD				RIOD		<u> </u>		
IV. SOURCES OF INFORMAT	TION (Cite specific ref	erences.	e.g., sti	ace files, sample an	alysis, reports)			

NUS FIT 3. Site inspection TDD No. F3-8810-15, December 6, 1989.



EDA

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT

I. IDENTIFICATION

EPA	PART9	- GE	NERATOR/TR	ANSPORTER INFORMATION	DC DC		OTE NUMBER	
N. ON-SITE GENERATOR								
01 NAME Washington	- 8 NUMBER							
03 STREET ADDRESS (P.O. Box. NFO P. exc.) 2215 Adams Place Northeast			04 SIC CODE					
oscity Washington	06 STATE	4	P CODE 10018					
III. OFF-SITE GENERATOR(S)	1 00		10010					
01 NAME		02 D	+ S NUMBER	01 NAME		02 D	+ 8 NUMBER	
03 STREET ADDRESS (P.O. Box, RFD P, etc.)			04 SIC CODE	03 STREET ADDRESS (P.O. BOX, RFD #, etc.)			04 SIC CODE	
05 CITY	06 STATE	07 ZI	P CODE	05.CITY	06 STATE	07 ZI	P CODE	
03 NAME		02 D	+ 8 NUMBER	Ot NAME		02 0	+ 8 NUMBER	
03 STREET AQORESS (P.O. Box, RFD #, etc.)	- 		04 SIC CODE	03 STREET ADDRESS (P.O. BOX, RFO #, etc.)			04 SIC CODE	
05 CITY	06 STATE	07 Z	PCODE	05 CITY	06 STATE	07 ZIP COOE		
IV. TRANSPORTER(S)								
01 NAME	······································	02 O	+ 8 NUMBER	OI NAME N/A			02 D + B NUMBER	
03 STREET ADDRESS (P.O. Box, RFD #, etc.)			04 SIC CODE	03 STREET ADDRESS (P.O. Box, RFD #, etc.)			04 SIC CODE	
05 CITY	06 STATE	07 ZI	P CODE	05 CITY	06 STATE	Q7 Z1	PCODE	
O1 NAME		02 D	+ 8 NUMBER	01 NAME N/A			02 D + 8 NUMBER	
03 STREET ADDRESS (P.O. Box, RFO ♠, etc.)			04 SIC CODE	03 STREET ADDRESS (P.O. Box, RFD #, etc.)			04 SIC CODE	
05 CITY .	06 STATE	07 Z	IP CODE	OS CITY OS STATE		07 ZIP CODE		
V. SOURCES OF INFORMATION (Cite spec	fic references.	e.g., st	ate files, sample anal	ysis, reports)	<u></u>			
NUS FIT 3. Site inspection	. TDD No	o. F:	3-8810-15, De	cember 6, 1988.				



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 10 - PAST RESPONSE ACTIVITIES

. IDENTIFIC	ATION
STATE	02 SITE NUMBER

				007
II. PAST RESPONSE ACTIVITIES				
01 D a. WATER SUPPLY CLOSED 04 DESCRIPTION	02	DATE0	3	AGENCY
None reported or observed				
01 8. TEMPORARY WATER SUPPLY PROVIDED 04. DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 C SERMANENT WATER SUPPLY PROVIDED 04 DESCRIPTION None reported or observed	02	DATE0	3	AGENCY
01 D SPILLED MATERIAL REMOVED 04 DESCRIPTION	02	DATE0	3	AGENCY
None reported or observed				
01 ☐ E CONTAMINATED SOIL REMOVED 04 DESCRIPTION	02	OATE 0	3	AGENCY
None reported or observed				
01 D F WASTE REPACKAGED 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 G. WASTE DISPOSED ELSEWHERE 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 HON-SITE BURIAL 04 DESCRIPTION	02	DATE0	3	AGENCY
None reported or observed				
01 ☐ I.IN SITU CHEMICAL TREATMENT 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01	02	DATE 0	3	AGENCY
01 K IN SITU PHYSICAL TREATMENT 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 LENCAPSULATION 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 M. EMERGENCY WASTE TREATMENT 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 O N. CUTOFF WALLS 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 ☐ O EMERGENCY DIKING/SURFACE WATER DIVERSION 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				
01 ☐ P CUTOFF TRENCHES/SUMP 04 DESCRIPTION None reported or observed	02	DATE0	3	AGENCY
01 🖾 O. SUBSURFACE CUTOFF WALL 04 DESCRIPTION	02	DATE 0	3	AGENCY
None reported or observed				

3	EP/

POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 10 - PAST RESPONSE ACTIVITIES

I. IDENTIFICATION

01 STATE 02 SITE NUMBER
007

H. PAST RESPONSE ACTIVITIES (Continued)			
01 C R. BARRIER WALLS CONSTRUCTED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 🗆 5. CAPPING/COVERING 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 T BULK TANKAGE REPAIRED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed.			
01 U GROUT CURTAIN CONSTRUCTED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 U 80TTOM SEALED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 U. W. GAS CONTROL 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 U x FIRE CONTROL 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 ☐ Y. LEACHATE TREATMENT 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 ☐ Z. AREA EVACUATED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			
01 1 ACCESS TO SITE RESTRICTED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or <u>observed</u>			
01 ☐ 2. POPULATION RELOCATED 04 DESCRIPTION	02 DATE	03	AGENCY
None reported or observed			_
01 G 3. OTHER REMEDIAL ACTIVITIES	02 DATE	03	AGENCY

HI. SOURCES OF INFORMATION (Cite specific references, e.g., state files, sample analysis, reports)

NUS FIT 3. Site inspection TDD No. F3-8810-15, December 6, 1988.



POTENTIAL HAZARDOUS WASTE SITE SITE INSPECTION REPORT PART 11 - ENFORCEMENT INFORMATION

I. IDENTIFICATION 02 SITE NUMBER 007 01 STATE DC

. ENFORCEME	NT INFORMATION
)1 PAST REGUL	ATORY/ENFORCEMENT ACTION TYPES NO
2 DESCRIPTIO	N OF FEDERAL, STATE, LOCAL REGULATORY/ENFORCEMENT ACTION
	ct of Columbia, Government Environmental Control Division, Hazardous Waste Section, sampled und the site in March, April, and June 1986.
No other re	egulatory or enforcement action has taken place at this site.
	\cdot
L SOURCES O	FINFORMATION (Cite specific references, e.g., state files, sample analysis, reports)
34444230	The second of th
NUS FIT 3.	Site inspection TDD No. F3-8810-15, December 6, 1988.

SECTION 6

TDD No.: F3-8810-15



6.0 REFERENCES FOR SECTIONS 1.0 THROUGH 5.0

- United States Geological Survey. Washington West, D.C. Maryland Virginia Quadrangle,
 7.5 Minute Series. <u>Topographic Map</u>. 1965, photorevised 1983. Combined with Washington East, D.C. Maryland Quadrangle, 7.5 Minute Series. <u>Topographic Map</u>. 1965, photorevised 1979; Alexandria, Virginia District of Columbia Maryland Quadrangle, 7.5 Minute Series. <u>Topographic Map</u>. 1965, photorevised 1983; and Anacostia, Maryland District of Columbia Quadrangle, 7.5 Minute Series. <u>Topographic Map</u>. 1965.
- 2. NUS Corporation, FIT 3. Site inspection; site visit. TDD No. F3-8810-15, December 6, 1988.
- Milosevic, Milan, Washington Plating Company, with Paul Persing, NUS FIT 3. Meeting. April 9, 1987.
- 4. Milosevic, Milan, Washington Plating Company, with Elizabeth Coughlin, NUS FIT 3. Telecon. March 27, 1989.
- 5. Milosevic, Milan, of Washington Plating Company, with Elizabeth Coughlin, NUS FIT 3. Telecon. March 30, 1989.
- 6. Bacon, Byron, District of Columbia, Department of Consumer and Regulatory Affairs, with Paul Persing, NUS FIT 3. Meeting. April 9, 1987.
- 7. Bacon, Byron, District of Columbia, Department of Consumer and Regulatory Affairs, to Angelo Tompros, Chief, Pesticides and Hazardous Waste Management. Correspondence. March 1986.
- 8. Bacon, Byron, District of Columbia, Department of Consumer and Regulatory Affairs, to Angelo Tompros, Chief, Pesticides and Hazardous Waste Management. Correspondence. February 1986.
- 9. Collier, James, Environmental Control, with Paul Persing, NUS FIT 3. Telecon. March 24, 1987.
- District of Columbia, Department of Public Works. Sewer Map. February 21, 1980.

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- 11. Kessler, Richard, Washington Suburban Sanitary Authority, with Paul Dietrich, NUS FIT 3. Telecon. March 17, 1987.
- 12. Fallin, Wayne, Washington Suburban Sanitary Authority, with Gilbert Marshal, NUS FIT 3. Telecon. October 21, 1985.
- 13. Cameron, Craig, Fairfax County Water Authority, with David D. Doran, NUS FIT 3. Telecon. September 21, 1986.
- 14. Eunpu, Floyd, Fairfax County Water Authority, with Edward Jamison, NUS FIT 3. Telecon. April 3, 1989.
- 15. Johnston, P.M., United States Geological Survey. Geology and Groundwater Resources of Washington, D. C. and Vicinity. Geological Survey Water Supply Paper, 1776, 1976.
- Mack, F.K., Maryland Geological Survey. Groundwater in Prince Georges County. Bulletin 29, 1966.
- 17. National Oceanic and Atmospheric Administration. <u>Climatography of the United States.</u>
 Local Climatological Data. Annual Summary with Comparative Data. Laural, Maryland. 1980.
- 18. United States Department of Commerce, National Climatic Center. <u>Climatic Atlas of the United States.</u> 1979.
- United States Department of Commerce, United States Printing Office. Rainfall Frequency Atlas of the United States. Technical Paper No. 40, 1963.
- Mr. O'Brian, United States Population Census Bureau, with Paul Persing, NUS FIT 3. Telecon.
 May 5, 1987.
- 21. Wolflin, John P., United States Department of the Interior, Fish and Wildlife Service, Division of Ecological Services, to Garth Glenn, NUS FIT 3. Correspondence. February 7, 1989.

TDD No.: F3-8810-15

22. Government of the District of Columbia, Department of Public Works Water and Sewer Utility Administration. Wastewater Discharge Permit Applications. August 11, 1986.

23. Donnelly, Daniel K., Annapolis Laboratory of EPA, to Neilima Sengalia, District of Columbia Hazardous Waste Section. Correspondence. May 12, 1986.

SECTION 7

TDD No.: <u>F3-8810-15</u>



7.0 LABORATORY DATA

7.1 Sample Data Summary

The attached data summary contains only compounds which were identified as detected in at least one sample. The complete list of compounds analyzed for, their results, and the associated detection limits are located as an appendix. Results for tentatively identified compounds appear following the organic data section of this report.

The following codes are used in the data summary to indicate the confidence in the laboratory results:

CODES RELATING TO IDENTIFICATION

(confidence concerning presence or absence of compounds):

U = Not detected. The associated number indicates approximate sample concentration necessary to be detected.

(NO CODE) = Confirmed identification.

B = Not detected substantially above the level reported in laboratory or field blanks.

R = Unreliable result. Analyte may or may not be present in the sample. Supporting data necessary to confirm result.

N = Tentative identification. Consider present. Special methods may be needed to confirm its presence or absence in future sampling efforts.

CODES RELATED TO QUANTITATION

(can be used for both positive results and sample quantitation limits):

J = Analyte present. Reported value may not be accurate or precise.

K = Analyte present. Reported value may be biased high. Actual value is expected to be lower.

E = Analyte present. Reported value may be biased low. Actual value is expected to be higher.

UJ = Not detected, quantitation limit may be inaccurate or imprecise.

UL = Not detected, quantitation limit is probably higher.

OTHER CODES

Q = No analytical result.

SAMPLING DATE(s): 12-6-1 STATE / COMP to CASS. 5/1-11 Washington Plating TOD WUMBER: F3-8810-15 CASE NUMBER: 11032 FEA WIMEST: LAB MAMES: IT Fitt(org), JTC Environmental Consultants (inorg) 64076 (19a77 53.078 SAMPLE NUMBER: CY071 CY069 CY070 SAMPLE ID: 5-1 9-2 9-3 9-4 1 -14-1 2524 An-flank Eletari, */200000 - ---LOCATION: Off-site off-site off-site 544-211E - 3-4 -- 2.4-1.2112 accessbackyd resdotaccess rechnistrate out access concrete concrete concrete comprete 10 Biressyet 35 (56) drk loamy drk.rchloamy 15989 sinapydayd dif sandblormarb e hear sandy.pebbly 7.3 6.5 6.2 5, 1 . 3 5.7 FIELD MEASUREMENTS: CE 75 PERCENT SOLIDS: 98.72 84.6% 84.4 TYPE OF DATA: ***** VOLATILES DILUTION FACTOR: 1.0 1.2 1.2 . . . 1.3 64074 19677 197.78 DET. LIMIT SAMPLE NUMBER: **EY069** CY070 CY071 Fx677 $\subseteq \phi_{i}(\tau)^{\frac{1}{2}}$ 67473 £7075 CROL (*=IDL) ug/kg ug/kg. 13,4 14.00 B 41,06.9 5.00 esthylene chioride 15.00 P 11.70 9 10.00 acetone 5.00 carbon disulfide 5.00 tetrachloroethene 1.00 B 1.65 \$ 3.00 B 1.40 B 5.00 1,1,2,2-tetrachloroethane 5.00_taluene $1.86 \pm$ 5.00 ethylaenzene 1 inv 5 2.00 € 5.00 styrene TYPE OF DATA: ****** 43.0 41.0 42.0 14.0 44.0 97.0 1.0 1,5 DILUTION FACTOR: 34.0 57171 92.7 74079 10.014 0:075 BET. LIMIT SAMPLE NIMBER: CYCAS €₹976 51671 es iks 99/89 __10.00_1.3-dichlorobenzene 14 (sp. 1 50:00 bearaic acid 100.00 J 280.09J* 10.00 2 10.00 10.00 10.00 20.00 10.0 190,00 1 87.00 3 1700.00 L 250.00 3 120.09 L 35,00 J 53,00 1 *70.00 10.00 acenaphthylene 51.00 1 37.0() 259,00 3 11 35,99 3 150. 10.00 acenaphthene \$40,000 10.00 dibenzofuran 160.60 110,00 J 400,60 3 740,00 [171,507 10.00 fluorene 200,903* 210,00 3 690.00 120,04 1 10.00 N-mitrosodiphenylamine 50.00 pentachlorophenoi 120,00 P 92.00 8 Fait 10 10.00 phenanthrene 2982.00 400,00 4359.00 1421/02 420.003* 15: (4) 1 490.00 1100.00 10.00 anthracene .ga nilis Iloyek ⊩ 220.00 P 77.95 8 27. July 6 210.008* 10.00 di-m-butyl obthalate 775,38 5100,00 (4800.00 3900,00 10.00 fluoranthere 2400.00 7700.00.t. 10.00 pyrene 4300.00 8 6500.00 J 1170 00 9 12000.00J# 1300.00 B 410.00 B 10.00 butylbenzyl obthalate 1700,00 1709.00 05.00.00 10.00 befizoralenthracene [500.]].P Trife You 9 720.09 5 10.00 bis(2-ethylbevyl) onthalats 1800.00 1496.90 10.00 chrysene 170.001* 10.00 di-m-octyl ohthelate 217 55 1400,00 # 10.00 benzo(b)fluoranthene 2500.00 1700.50]* 1600,00 10.00 benzo(k)fluoranthene 2400.00 read to the second - 7 × 10.00 benzolatovnese erspires 10.00 indens[1,2,3-od]pyrens 24,55 % 100.00 J 10.00 dibenz(a,h)anthracene 10.00 benzo(q,h,i)perviene 759,00 1400.00 210,00 450 3 TYPE OF DATA: ****** PESTICIDES

1700.0

DILUTION FACTOR:

1400.0

SITE MAME: TOO NUMBER:	Washington Plating F3-8810-15		CASE	ING DATE(s): 1 NUMBER:11032		STATE/COUNTY COME SPA NUMBES:	Fr 05 -17 55 65 7		(
LAB NAMES:	IT Pitt(org),JTC Envir	onmental Consul	tants (inorg)	* * * * * *								
	SAMPLE NUMBER:	EY069	EY070	CY071	0.6072	C4073	[YA74	57075	€r97 5	Ev977	31078	
	SAMPLE ID:	5-1	9-2	9-3	9-4	9-5	S-Fart	Sd-1	34-1	An-tlank	Sablank	
	LSCATION;	Off-site	off-site	off-site	0++-5155	tit te S	oxxxxx nativo	tesement	tagorar.	o.ari	ካ (a	
		pub access	accessbackyd	resdotaccess	resdotaties	=	nut n ancesa	ar -şifə	25-5116			
		concrete	concrete	concrete	contrete		no Romeseveç	Eu-1	61 3666			
		sandy.pebbly	drk loamv	drk.rchloamy	lpa=v		lear-Rawdus*	eanoblekgerb	rlear			
	PH:	7.3	6.5	6.2	5.7	*	7.0	7.7	5, 3			
	FIELD MEASUREMENTS:						0.1 pp=					
	PERCENT SOLIDS:	98.72	84,5%	84.5%	7E.2'.	50	7	75.75				
TYPE OF DAT	A: ***** PESTICIDES	*******	**********	*********	**********		***********	**********	***********			***************
	DILUTION FACTOR:	1300.0	1760.0	1600.0	1699.4	:70 - 3	175.	470.3	1.0	ls.		
BET. LIMIT	SAMPLE NUMBER:	CY069	CY070	CY071	CY072	CY 073	eyaze	£¥975	[٧٨74	54977	CY078	
CROL_{**IDL) UNITS:	_ug/kg	ug/kg	ug/kg	ug/kg	uq/kg	9g/Fg	EQ7-EQ		un/1	ña/	
1.00 d	ieldrin				510,0,8		94m, jr. 5					
	.4'-DOT	380.00 P	₿ộბ.gọ ƙ	749.00 R	1996,6436							
0.05 a	roclor-1248				98 99.06 0							
` Fassen	te: #####################	*************	************					4444				

C= confirmed by 60/MS

data validated by RC
*= reported from dilution or re-extract

7 F 6 6	· Washington Plating Re F3-8810-15	i I	CASE	[ING DATE(s): 1 NUMBER:11032	1		: 05-11 05067	1	1 ([1 1	1		
LAR MAKES	: IT Pitt(org),JTC Envir	ormental Consul	tants (inorg)											
: 1	SAMPLE NUMBER:	MC4048	MCY069	MCY070	M[V[7]	MC+372	MCY(73	MEY974	MCV075	MCY074				-
<u> </u>	SAMPLE ID:	5-1	E-2	5-3	<u> 5</u> - 4	9-5	9-590	5d-1	54-1	4q-51acx				
- 1 . 1	LOCATION:	Off-site	off-site	off-site	off-site	515 €4 S-1	iş+-eşteştê	gasement	545 998 61	ճ[գո⊧				
1		Dub access	accessbackvo	resdntaccess			topic access	्र- ११ रेट	98-31 te					
****		concrete	concrete	concrete	concrete		no stresskeo	5W-1	ñ5 ⊍5 P 9					
1	PH:	sandy,pebbly	dr∮ lo∋my	drk,rchloamy	loamv 5.7	3.3	loamysandus*	eandblokgarb	clear , -					
1	FIELD MEASUREMENTS:		6.5	6.7	٠. ١	7, 2	0.3 000	7,2	6.7					
	PERCENT SOLIDS:	98.2%	81.0%	84.67	78.2%	99,30	73.1%	55.7%					•	
TYPE DE D	ATA: ###### INOPGANICS		0:,V4 :**********							***********	**********	***********	**********	********
in the second of	DILUTION FACTOR: : GFA		0.240	0.240	0.260	(.209	0.170	0.360	1,000	1.900				
	; ICP		0.240	0.740	0.260	0.269	9,770	0.360	1,000	1,000				
·	: Hg	0.510	0.620	0.590	9.649	0.519	0.689	0.900	1.000	1.000				
	: CN	0.200	0,250	0.240	9.260	0.200	0.270	0.360	1,000	1.900				
			MENALE			45.13.43			W0075	450437				
DET. LIMI		MCY068	MEY069	MCY070	MCY071	MCY072	hCY073	MCY074	MCY975	#CY076				
CROL(#=]		<u>mg/kg</u> 2710.00 L	#g/kg 5130,00 L	#g/kg 6450.00 L	<u></u>	<u>59/kg</u> 2549,00 L	#9/kg 4460,00 L	<u>09/kg</u> 4440,00 L	ug/l	49/1				
\$ 4 164.00 * 48.00		21.60	2120'00 F	6450.00 F	2910*****	£290.00 L	4447 (0.) 7	75, 4 0						
# 5.00	,	5.80 L	5.80 8	7.70 J	12,36	6,10	10.20_1	20.60	5.40 B					
8.60		7520.00	390,00	559.00	475,00	2176 46	704,00	1(200,00	(45.00	-				
	bervliium			0.77	1.20	_		,						
4.30		4,00	1.60	1.90	5.19	5,00	13.50	13.40						
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SAMPLE DATA SUMMARY

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3) SITE NAME

4) STATE & COUNTY CODE 5) ORGANIC OR INCRGANIC 6) LABORATORY NAME 1 11) DILUTION FACTOR

7) TRAFFIC REPORT #

8) SAMPLE IDENTIFIER

10) CONCENTRATION

12) PH 13) FIELD MEASUREMENTS

14) PERCENT SOUD

4320C-1 and 4230C-5 are field duplicates



TDD No.: F3-8810-15

ORIGINAL (Red)

7.2 Quality Assurance Review

7.2.1 Organic Data: Lab Case 11032

7.2.1.1 <u>Introduction</u>

Seven solid and two aqueous samples were analyzed for acid, base-neutral, and pesticide/polychlorinated biphenyl (PCB) compounds through the EPA Contract Laboratory Program (CLP). Included in the sample set were one field duplicate pair and one field blank. Also included was one aqueous blank to be analyzed for volatile organic compounds only.

The data have been fully reviewed to determine the usability of results according to the National and Regional Guidelines. (Areas examined in detail are listed in the Support Documentation appendix.) Data quality was acceptable for most compounds, with detection limit capability demonstrated by meeting criteria for holding times, surrogate and matrix spike recoveries, and instrument tuning and calibration. Detection limit capability is questionable for some acid compounds in a few samples. Blank contamination affected low levels of most volatile compounds and a few semivolatile compounds.

Principal areas of concern include blank contamination and low acid surrogate recoveries for one sample in particular.

7.2.1.2 Qualifiers

- All results for methylene chloride, acetone, 1,1,2,2-tetrachloroethane, tetrachloroethene, toluene, ethylbenzene, styrene, xylenes, pentachlorophenol, and di-n-butyl phthalate have been flagged as undetected due to blank contamination (B). All results for these laboratory contaminants are not significantly higher than the levels detected in all associated blanks. Other results that are flagged (B), due to blank contamination, include butylbenzyl phthalate in samples CY070, CY071, CY071, CY074, and CY075 and bis(2-ethylhexyl) phthalate in samples CY070, CY071, CY072, CY073, CY074, and CY076.
- The volatile fraction of sample CY069 was analyzed immediately after the third laboratory blank, which was run right after the daily calibration standard. Both the blank and this sample contained low levels (1 to 6 ug/l) of many later-eluting compounds. The failure of the gas chromatographic system to completely purge the trap of all compounds may be a contributing factor for this observed carryover.

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DRITINAL (Red)

• The laboratory diluted and re-analyzed the semivolatile fractions of solid samples CY069 and CY073 (field duplicates) because of high levels of butylbenzyl phthalate. With the exception of pyrene and di-n-butyl phthalate, the reviewer has reported the highest levels of all compounds detected in both analyses. Specifically, for CY069, all results were reported from the initial analysis except for fluorene, anthracene, di-n-octyl phthalate, benzo(b)fluoranthene, benzo(k)fluoranthene, and benzo(a)pyrene. For CY073, butylbenzyl phthalate, benzo(a)pyrene, and benz(a)anthracene have been reported from the re-analysis.

- Butylbenzyl phthalate was calculated manually by the reviewer for the dilution of CY069; this
 compound was not identified as a target compound by the laboratory software but was
 present as a tentatively identified compound (TIC) in the re-analysis of the sample. The result
 has been flagged as estimated (J) because of errors inherent in the manual calculation.
- The result for dibenz(a,h)anthracene has been flagged as tentatively identified in sample CY069. The result was very low, and the sample spectrum exhibited only a marginal match with the reference spectrum. A high hydrocarbon/siloxane background made identification difficult, but the presence of many other related polynuclear aromatic hydrocarbons (PAHs) corroborate the presence of dibenz(a,h)anthracene.
- The laboratory re-extracted and re-analyzed the semivolatile fraction of sample CY075 because of several low surrogate recoveries. The recoveries for 2-fluorobiphenyl and terphenyl were similarly low in both analyses, and the other surrogates displayed very similar recoveries as well. The reviewer has reported the highest levels of all compounds detected, with the exception of compounds considered attributable to blank contamination (see the following table). Benzoic acid, indeno(1,2,3-cd)pyrene, dibenz(a,h)anthracene, and benzo(g,h,i)perylene were the only non-artifact compounds reported from the re-analysis of this sample.
- Several results have been flagged because of blank contamination, while others have been
 reported as either unreliable or confident for samples CY069, CY073, and CY075, even though
 these compounds were detected in the blanks. The following table itemizes which phthalate,
 pentachlorophenol, or N-nitrosodiphenylamine results were reported for these two samples
 and the reasons behind the decisions.

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Sample	Compound	Result (ug/kg)	Analysis	Decision Criterion
CY069	pentachlorophenol	120	initial	1
	di-n-butyl phthalate	not detected	dilution	2
	butylbenzyl phthalate	12,000 ug/kg	dilution	4
	bis(2-ethylhexyl) phthalate	3,500	initial	3
CY073	N-nitrosodiphenylamine	not detected	initial	2
	di-n-butyl phthalate	1,500	initial	1
	butylbenzyl phthalate	30,000	dilution	4
	pyrene	not detected	initial	2
	bis(2-ethylhexyl) phthalate	2,100	initial	1
	pentachlorophenol	73	initial	6
CY075	N-nitrosodiphenylamine	not detected	initial	2
	di-n-butyl phthalate	210	re-analysis	1
	bis(2-ethylhexyl) phthalate	23,000	initial	5
	butylbenzyl phthalate	270	re-analysis	1

<u>Decision Criteria</u>

- 1. Both results were questioned by blanks; the lowest value was reported and flagged (B).
- 2. One result was questioned by blanks; the other result was not detected. The not-detected result was reported.
- 3. One result was greater than 10 times but less than 20 times the highest blank value; the other result was questioned by the blanks. The higher result was reported and flagged as unreliable (R). Further information is necessary to verify the presence of this compound at this location.
- 4. Both results were at least 20 times the highest blank result. The highest result was reported and is considered confident. (Other high levels of phthalates may corroborate the presence of this compound at this location for sample CY069.)

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5. One result was greater than 20 times the highest blank, but the second result was in the same range as the blanks. The highest result was reported and flagged (J) due to disagreement between the two analyses. However, further information may be useful in verifying the presence of this compound at this location.

- 6. The initial result was questioned by the blanks. This result was reported because the detection limit for the diluted re-analysis was too high to enable detection of this compound at the level seen in the undiluted analysis.
 - Surrogate recoveries were low for two base-neutral extractable compounds for solid sample CY075. This indicates that detection limits for many undetected polyaromatic compounds may be higher than reported in this sample, and results for the PAHs that were detected may be biased low and have been flagged (L). In addition, surrogate recoveries for nitrobenzene and tribromophenol were slightly low, although contractually acceptable. Detection limits for some substituted aromatic compounds, as well as for highly substituted phenols, may be slightly higher than reported in this sample.
 - Slightly low surrogate recoveries for tribromophenol were also observed for solid samples CY069 and CY072. Detection limits for highly substituted phenols may be slightly higher than reported in these two samples.
 - The recoveries for d₅-phenol were low for all aqueous samples, including the laboratory blanks. The matrix spike/matrix spike duplicate recoveries for phenol and 4-nitrophenol were low for aqueous sample CY076 as well. This suggests a problem with the extraction of phenol and possibly other phenolic compounds from an aqueous matrix. The detection limits for phenol and 4-nitrophenol in particular may be higher than reported in all aqueous samples.
 - Even though unusual, the result for 1,3-dichlorobenzene is considered confident in sample CY069. The sample spectrum matches the reference spectrum well, and the compound eluted at the expected retention time. However, this compound was not detected in the field duplicate, sample CY073, possibly due to the very low instrument levels involved. (Also, the concentrations of most compounds were lower in CY073 than in CY069, which could also account for the absence of this compound in CY073.)

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There was generally good precision for the results between field duplicate samples CY069 and CY073. However, imprecision was seen for 1,3-dichlorobenzene, acenaphthene, pyrene, butylbenzyl phthalate, di-n-octyl phthalate, and indeno(1,2,3-cd)pyrene. In most cases, the compound was detected in only one sample. Results for these compounds have been flagged as estimated in samples CY069 through CY074, unless previously flagged (B).

- The result for benzo(a)pyrene is considered estimated in sample CY073. The initial, undiluted result was very low (only 0.34 ug/l), instrument level, whereas the diluted instrument level result was 2.7 ug/l. Sample inhomogeneity may be the cause of the observed difference.
- All results for dieldrin and 4,4'-DDT (except DDT in sample CY072) have been flagged as unreliable (R). These pesticides are represented by a single chromatographic peak, and the presence of interferences eluting within the expected retention time window of these compounds can cause false positive results. Without the presence of related breakdown or parent compounds, further information would be necessary to verify the presence of these compounds at these locations. The presence of relatively high levels of PAHs and phthalates in many samples may have contributed to these interferences.
- Detection limits may be higher than reported for DDT and dieldrin in all solid samples. The
 high levels of PAHs in sample CY070 may have enhanced the peaks representing these two
 matrix spike compounds, resulting in the high recoveries seen for this sample. Because
 similar levels of PAHs exist in all solid samples, detection limits may be affected in all solid
 samples.
- The results for Aroclor 1248 and DDT are considered confident in sample CY072. Both compounds were confirmed by gas chromatography/mass spectrometry (GC/MS), and the Aroclor displayed good peak pattern matching quality and area ratios with the reference standards on both chromatographic columns. Because of the presence of multi-peak PCBs in this sample, compounds related to DDT (i.e., DDD and DDE) could not be distinguished. The peaks representing the Aroclor may have masked the single peaks representing DDD and DDE.

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ORIGINAL (Red)

Detection limits for 2-butanone may be higher than reported in all samples. The continuing
calibration response factors were less than 0.05. This is a common problem with this
compound because EPA requires the use of a quantitation ion that is a minor component of
the mass spectrum for this compound.

- Tentatively identified compounds that are not demonstrated artifacts or laboratory contaminants are summarized immediately following this report.
- Sample results that are below the calibration range of the analysis have been flagged as estimated (J) on the data summary, where no other flag exists.

7.2.1.3 Support Data

The Support Documentation appendix to this report documents the above findings associated with blank contamination, the flagging of one result as tentative, the manual calculation of one result by the reviewer, low semivolatile surrogate and matrix spike recoveries, and the low response factors for 2-butanone. (Issues pertaining to laboratory contractual compliance are found on a separate summary directed to the laboratory deputy project officer.)



s	AMPLE DATA S	SUMMARY	: ORGANIC TENTATIVELY IDENTIFIED COMPOUNDS
SAMPLE ANALYSIS NUMBER FRACTION (VOA/BNA)	S ESTIMATED N CONCENTRATION	QUALIFIER CODE	
CYOG9 VOA	3100 rg/kg 270 1 420	10T 10T 150	Carboxylic acids (5) aidehycles (2) 9-H Carbazole Cizitan Cistiz PAH-2-methylanthracene
1 3NA	1600 J	150	CICHIZ PAH- Such as 2-methyl anthracene 9,10-anthracene dione C14 H& D2
CYOTI VOA BNA		150 150 150 150	CIHIO PAH-SUCHAS I-methy/maphthalene CIZHIZPAH-SUCHAS di methy/naphthalene CISHIZPAH-SUCHAS 4-methy/phenenthrene CIGHIZPAH-SUCHAS 2-pheny/maphthalene
240TZ VOA	1300 139 149 1550 1550 1	150	Trichloro Suphenyl Tetra Chloro bi phenyl 4-4-100T CISTARPAH SUCH as 2-methylanthracene.
Cyo-B VOA BNA	1 200 M3/ka 1 3 90 1 1 2 30 1 1 5 30 1 1 2 400 1	9 150 UNK	Cerboxylic a wid galactifol denivative Cistlic PAH such as 2-methyl anthrocene anthrocenediane Ciutiala satd NC Unknown
9074 VOA		1	canboxylic acid galactital derivative Canboxylic acid plus 2-methylanythracene anthracenedione Sold NC

SUS = <u>SUSPECTED FALSE POSITIVE RESULT:</u> Compound is either a common laboratory contaminant, or else a possible reaction byproduct (artifact) attributable to the chemical reagents used for sample preparation and analysis. This result is suspect even though this compound was not found in any associated blanks.

DEFINITIONS OF QUALIFIER CODES

UNK = UNKNOWN COMPOUND: Library search result unreasonable or of very low matching quanty.

TOT = TOTAL CONCENTRATION REPORTED: Represents the sum of several compounds detected all belonging to the same chemical class.

ISO = OR ISOMER: Compound identification is not selective for this isomer only. This result may instead represent the presence of a similar compound comprised of the same atoms bonded together in a different arrangement or substitution pattern.

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CYOTO VOA	(VOA/BNA) VALUE UNITS		ND = none detected X= number of compared harcategory
SNA 610 19 150 (12 Hz PAN-Such as d)methy naphthetere 2600 Galactital clenuative 23000 Carboxylic Ocid 500 SSO C17 Hz PAN Such as H-methyl pyrene 420 henz (d, e) anthracen-7-one C17 H100 450 UNK poss answetic or PAH 5800 TOT UNK poss answetic (2) V 2100 UNK poss answetic (2)	C4075 1VOA 1		
23000 Carboxylic acid 500 Iso Cithic phil sychan H-methyl pyrene 420 heng (d. e.) anthracen-7-one CithioO 450 unk poss anomatic or phth 5800 Tollunk (Inknowns (2) 1000 Unkleit Unknowns (2) Cyore VOA 1000 UNK	7 7 N 10 1 10 10 10 10 10 10 10 10 10 10 10 1	150	
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DEFINITIONS OF QUALIFIER CODES:

SUS = SUSPECTED FALSE POSITIVE RESULT: Compound is either a common laboratory contaminant, or else a possible reaction byproduct (artifact) attributable to the chamical reagents used for sample preparation and analysis. This result is suspect even though this compound was not found in any associated planks.

UNK = UNKNOWN COMPOUND: Library search result unreasonable or of very low matching quality.

TOT = TOTAL CONCENTRATION REPORTED: Represents the sum of several compounds detected all belonging to the same chemical class.

ISO = OR ISOMER: Compound identification is not selective for this isomer only. This result may instead represent the presence of a similar compound comprised of the same atoms bonded together in a different arrangement or substitution pattern.

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ORIGINAL (Red)

7.2.2 Inorganic Data: Lab Case 11032

7.2.2.1 <u>Summary</u>

Seven solid and two aqueous samples were analyzed for total metals and cyanide through the EPA

CLP. Included in the sample set were one duplicate pair and one field blank.

The data have been fully reviewed to determine the usability of results according to the National and Regional Guidelines. (Areas examined in detail are listed in the Support Documentation appendix.) Data quality was good for most metals and for cyanide. Detection limit capability was demonstrated for most elements by meeting criteria for holding times, spike recoveries, calibration check standards, low-level standards, and linear-range analyses. Low levels of several metals were detected in the

laboratory and field blanks, and some qualitative problems affected some of the results.

Principal areas of concern include blank contamination, spectral interference of silver from iron, a few low matrix spikes, laboratory duplicate imprecision, one miscalculated result, and a few variant

post-digestion spike recoveries.

7.2.2.2 Qualifiers

• Based upon careful examination of data from this case and other CLP cases that employ the same operating parameters (namely wavelength), it has been determined that spectral interferences from iron caused substantial signal suppression for silver, resulting in negative instrument readings for all solid samples. The interfering wavelengths were determined from literature references. The sample data, as well as the interference check standard, displayed levels of iron approximately 2,700 times the level of silver (on an absolute value basis). The correlation coefficient for the iron/silver data from this case was 0.991. The detection limits for silver may be higher than reported in all samples except MCY076 and MCY075, which contained low levels of iron (under 1,000 ug/l). This is corroborated by a zero percent matrix spike recovery for silver in sample MCY069.

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Several metals were detected in the laboratory blanks. However, only results for two metals were affected. The results for arsenic in samples MCY069 and MCY075, as well as all positive results for cobalt except MCY074, have been flagged as undetected due to blank contamination (B). The levels of these metals in the associated samples are not significantly higher than the levels in all associated blanks.

- The result for barium was miscalculated by the laboratory for sample MCY074. The reviewer
 has reported the correct result, 11,200 mg/kg, on the data summary.
- The reviewer has reported 5.8 mg/kg for arsenic in sample MCY068. This metal was detected in two furnace analyses but was not detected in the method of standard additions (MSA) analysis. The possibility exists that the wrong sample was analyzed by MSA and that the positive result has been reported as a worst-case approach. In addition, this result has been flagged as biased low (L), as a result of slightly low post-digestion spike recoveries for both furnace analyses.
- The matrix spike recovery was 49 percent for arsenic in aqueous sample MCY075, and the post-digestion spike recovery was 70 percent for this sample. Blank contamination may be the cause for the low recovery [the duplicate result, 5 U (not detected), yields a recovery of 63 percent]. Since this is the only surface water sample in the case, only this one result may be slightly higher than reported. However, the result has already been flagged (B). A similar situation exists for thallium; the matrix spike recovery in this sample was 57 percent, and the post-digestion spike recovery was 91 percent. Because these data suggest digestion loss, the detection limit for thallium may be higher than reported for sample MCY075.
- Low matrix and post-digestion spike recoveries were observed for antimony in solid sample MCY069. This suggests sample-specific matrix suppression upon analysis for this metal. The detection limit may be higher than reported for antimony in this sample.
- The matrix spike recovery for selenium in sample MCY069 was reported from MSA data, which do not always reveal the cause for the low recoveries (matrix suppression versus digestion losses). The initial data reveal slightly low matrix spike recoveries for this sample and the laboratory duplicate (62 percent and 52 percent, respectively) and good post-digestion spike recoveries. This indicates digestion losses, and, therefore, all solid selenium results may be higher than reported and have been flagged (L). Detection limits may be higher than reported in all solid samples where selenium was not detected.

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ORIGINAL (Red)

The matrix spike recovery for silver was slightly low at 59 percent in aqueous sample MCY075.
 The detection limit may be higher than reported in this sample.

- The matrix spike recovery for cyanide was slightly high in solid sample MCY069. All solid results can be further considered estimated and may be slightly lower than reported.
- Slightly low correlation coefficients were observed for selenium in samples MCY069 and MCY070, as well as for arsenic in sample MCY073. These results may be considered estimated and have been flagged (J), where no other flag exists.
- Laboratory duplicate imprecision was observed for iron in aqueous sample MCY075 and for selenium in solid sample MCY069. All solid results for selenium may be considered further estimated, and the iron result in sample MCY075 has been flagged as estimated (J).
- All lead results were reported from the plasma data, except for sample MCY076 (the blank).
 Sample MCY075 was also analyzed by furnace, and excellent agreement was seen between the furnace and plasma data for this sample.
- The laboratory did not perform a serial dilution for the aqueous matrix, which consisted of one sample. Therefore, no observations regarding possible matrix influences on the data can be made.
- No cyanide was observed in sample MCY068. However, 64.6 mg/kg of this analyte was detected in the field duplicate, sample MCY072. No field duplicate imprecision was observed for any other analytes, suggesting that the wrong sample may have been analyzed for either one of the cyanide analyses. Careful examination of all cyanide data provided did not confirm this possibility, however. The result in sample MCY072 has been flagged as estimated (J), as have all other solid results for cyanide. Further information would be useful in verifying the presence of cyanide at the particular location represented by MCY072.

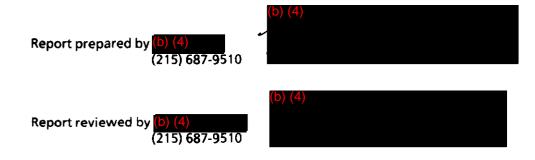
TDD No.: F3-8810-15



• The recoveries for the solid laboratory control sample (LCS) were low for aluminum, selenium, and silver. The recovery for silver was below EPA limits. High iron values in the LCS may account for the low silver recovery, and digestion loss may account for the low selenium recovery. According to the July 1987 Revision of the Inorganic CLP Statement of Work (SOW), "If the results for the solid LCS fall outside the control limits established by EPA, the analyses must be terminated, the problem corrected, and the <u>previous samples</u> associated with that LCS <u>re-digested</u> and <u>re-analyzed</u>."1 This was not done for silver for this case. The low recoveries for selenium and silver corroborate the low matrix spike recoveries discussed earlier in this report. Results for aluminum in all solid samples may be higher than reported and have been flagged (L).

7.2.2.3 Support Data

The Support Documentation appendix to this report documents the above findings associated with blank contamination, low spike recoveries, spectral interferences for silver, a miscalculated result, and laboratory duplicate imprecision. This report has been formatted to address those issues directly affecting the application of the data to the subject investigation. (Issues pertaining to laboratory contractual compliance are addressed on a separate form directed to the laboratory deputy project officer.)



¹Revision of Inorganic Statement of Work. July 1987. Section E, page 12.

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ORIGINAL (Red)

7.2.3 SAS 4320C - Hexavalent Chromium

Six solid samples were analyzed for hexavalent chromium by the Special Analytical Services (SAS) provision of the EPA CLP. Included in the sample set was one duplicate pair.

Analytical methodology was based upon method 3060 alkaline digestion for hexavalent chromium and method 7197 chelation/extraction flame atomic absorption for hexavalent chromium.

The data have been fully reviewed to determine the usability of results according to the National and Regional Guidelines. (Areas examined in detail are listed in the Support Documentation appendix.)

Data quality was good with respect to blank results, spike and duplicate performance, and instrument calibration. All recoveries and precision were well within quality control limits. No results were affected by any problems because there were no positive results and there were no problems observed for this analysis.

The Support Documentation appendix to this report includes blank, spike, duplicate, and calibration verification results.



SECTION 8

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8.0 TOXICOLOGICAL EVALUATION

8.1 Summary

Notable levels of several metals and cyanide were observed in water and sediment taken from the flooded basement of the plating room. However, the levels of inorganic contaminants, combined with the minimal amount of exposure expected, seem to indicate that no significant impacts should be expected.

Polycyclic aromatic hydrocarbons (PAHs) and bis(2-ethylhexyl) phthalate were detected in basement sediment at levels not expected to pose a significant hazard.

Soil samples obtained from the alley and backyards near the plating room revealed notable levels of lead, cyanide, barium, nickel, antimony, and cadmium. While lead levels were not necessarily atypical for urban environments, it is generally considered desirable to minimize all lead exposure. At the measured levels of other inorganic contaminants, no significant impacts are expected.

In one backyard soil sample, polychlorinated biphenyls (PCBs) were detected. Including tentatively identified PCBs, an EPA clean-up guideline for residential areas was slightly exceeded. Accidental ingestion of 100 mg of this soil would not be expected to result in significant impacts. Dermal absorption from soil is usually negligible, as PCBs are strongly soil bound.

In alley and/or backyard soil samples, 4,4'-DDT, PAHs, phthalates, and volatile organic compounds (VOCs) and semivolatile organic compounds (SVOCs) were detected. The reported soil levels of these contaminants are not expected to result in significant health effects.

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8.2 <u>Support Documentation</u>

8.2.1 Inorganic Contamination

The basement of the plating room was reported to be flooded with approximately four feet of water from backed-up wastewater and some storm water that could wash in through a hole in the wall. Some metal levels [chromium (113 ug/l), copper (46.6 ug/l), lead (1,290 ug/l), mercury (0.2 ug/l), nickel (1,810 ug/l), and zinc (328 ug/l)] were detected in excess of Ambient Water Quality Criteria (AWQCs), which are criteria used to judge typical surface waters. This water, however, is not typical surface water and is not expected to support aquatic life. Access to the room is restricted, except for a hole in the wall. Sediment samples were taken from the basement; this sediment was compared with typical nonpolluted eastern United States soils for the sake of perspective.² This sediment is not, however, a true soil but appears to consist of debris, washed-in dirt, and dirt that has settled out from the water. Antimony (76.4 mg/kg), barium (11,200 mg/kg), cadmium (13.4 mg/kg), chromium (2,900 mg/kg), copper (1,880 mg/kg), cyanide (10.6 mg/kg), lead (8,070 mg/kg), mercury (4.7 mg/kg), nickel (1,680 mg/kg), and zinc (14,500 mg/kg) were detected at levels above typical nonpolluted soil levels.² While occasional trespassing may provide occasional exposure opportunities, no significant adverse effects are expected.² Metals tend to adsorb onto particulates and would not be well absorbed in this situation.³ Metal toxicity is usually observed from high-level industrial exposure to dusts or fumes, which is not the situation at this site.4

Duplicate composite surface soil samples taken from alleys near the plating room revealed elevated levels of lead (up to 1,090 mg/kg), cyanide (64.6 mg/kg), barium (up to 2,520 mg/kg), nickel (up to 1,090 mg/kg), antimony (21.6 mg/kg), and cadmium (up to 5 mg/kg). Soil samples taken from backyards bordering the southern alley revealed elevated levels of lead (up to 1,660 mg/kg), cyanide (up to 9.1 mg/kg), cadmium (up to 5.1 mg/kg), and nickel (1,360 mg/kg).

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Lead was also detected in background soil at 502 mg/kg. The majority of lead compounds found in the urban environment result from leaded gas combustion.⁵ These types of lead are more heavily concentrated around roadways and garages; upper layers of roadside soil (within 25 meters of the road) may have as much as 2,000 ppm in excess of natural lead levels.⁵ On-site soil levels are not necessarily atypical for urban soils. However, because the site history involves the use of metals, the source of lead cannot be definitely determined. Lead has been seen to affect the hematopoietic, gastrointestinal, renal, and nervous systems.4 Accidental ingestion of lead-contaminated soil is usually a greater problem for children than adults; children are more sensitive to lead because of their developing nervous systems and greater lead absorption. 6 Lead, however, binds strongly to soil, decreasing its availability. Inadvertent ingestion of 100 mg of the most contaminated backyard soil would result in a lead intake of 166 ug. Dietary daily lead intake has been reported to be 119 to 274 ug per day for adults and 40 to 210 ug per day for children.⁷ A single exposure of 100 mg of the most contaminated soil would not be expected to produce significant health effects in and of itself. However, blood-lead increases of about 2 ug/dl per 1,000 mg/kg soil lead have been predicted after chronic lead exposure.8 This would represent a blood-lead increase of only about 3.3 ug/dl above baseline. As previously mentioned, urban residents may have an already high lead baseline. It is generally considered desirable to minimize all lead exposure, as people are exposed by a variety of sources, especially in the urban environment.

Cyanide has been reported to be nontoxic to humans at ingested levels up to 10 mg per day (5 mg per day for long-term consumption) because it is detoxified in the body.⁹ It is not considered to be an important environmental problem because of its low persistence and its metabolic biotransformation in the body.¹⁰ Accidental ingestion of 100 mg of the most contaminated soil would result in a cyanide intake well below the health-based risk reference dose (RfD).¹¹

Nickel, reported to be a possible site-related contaminant (see section 4.0), has been seen to cause dermatitis in sensitive individuals; toxicity has been observed after high-level industrial inhalation exposure.⁴ Nickel is also an essential element. Accidental ingestion of 100 mg of the most contaminated soil would not only result in a nickel intake (136 ug) below the RfD, but also below the adult daily dietary intake of 300 to 600 ug per day.^{6,11} Background soil nickel was detected at 717 mg/kg.

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OKIGINAL (Red)

Cadmium, barium, and antimony have produced toxicity at high levels in industrial settings.^{4,6} Like most metals, they are slowly absorbed and are not generally available when adsorbed onto soil particles. At on-site levels, no significant impacts are expected. The background cadmium level was reported to be 13.5 mg/kg, higher than any on-site soil samples.

8.2.2 Organic Contamination

No significant organic contaminants were detected in the water sample obtained from the basement.

PAHs were confidently and tentatively identified up to 33,830 ug/kg in basement sediment. PAHs are often found in the environment, occurring in coal and tar and forming from the incomplete combustion of organic material.⁶ At higher levels than those observed at this site, PAHs can cause dermatitis in sensitive individuals.⁴ PAHs adsorb strongly onto soil particulates, reducing their availability. As previously discussed, contact, if any, with this sediment is expected to be minimal. This water is not expected to support aquatic life.

DEHP was detected in basement sediment at 23,000 ug/kg. Phthalates, as plasticizers, are ubiquitous in the environment.⁶ It is interesting to note that the sediment sample was reported to contain garbage and debris. Phthalates are noted for their low acute toxicity and low chronic toxicity.⁶ DEHP is classified as a suspect human carcinogen, but the limited exposure potential to basement sediment makes a cancer risk impractical to quantitate.¹² No significant impacts are expected.

In the alley soil samples, PAHs were confidently and tentatively identified at levels up to 15,839 ug/kg. Phthalates, not including DEHP, were detected up to 30,000 ug/kg. 1,3-Dichlorobenzene (74 ug/kg) was also detected in these samples.

In soil taken from the backyards, PAHs were confidently and tentatively identified (up to 29,702 ug/kg), PCBs were confidently and tentatively identified (up to 10,650 ug/kg), and 4,4'-DDT was confidently and tentatively identified (up to 2,450 ug/kg).

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As previously mentioned, PAHs are practically ubiquitous in the environment.⁶ Background soil PAHs were detected up to 9,550 ug/kg. No significant non-carcinogenic effects are expected at on-site levels. Some PAHs have been classified as suspect human carcinogens: benz(a)anthracene, chrysene, benzo(b)fluoranthene, benzo(k)fluoranthene, benzo(a)pyrene, indeno(1,2,3-c,d)pyrene, and dibenz(a,h)anthracene.¹² Theoretical increase in cancer risk cannot be quantitated because PAH cancer potencies are being re-evaluated by EPA; due to the no-threshold theory of carcinogenicity, some increase in cancer risk cannot be ruled out.

The total level of PCBs slightly exceeded an EPA clean-up guideline of 10,000 ug/kg for residential or unlimited-access areas.¹³ These persistent, lipophilic compounds have been associated with chloracne and liver damage at high levels.⁴ They are suspect human carcinogens.¹² Generally, toxicity has been associated with large-scale industrial inhalation exposure or accidental ingestion of large amounts of PCBs.^{13,14} In contrast, exposure to PCBs at this site is likely to occur via dermal contact with or accidental ingestion of small quantities of soil or inhalation of dust or volatilizing PCBs. Dermal contact with PCBs is usually considered to be a relatively insignificant exposure route, especially from soil, except when oil is present to act as a vehicle.¹³ The samples at this site were described as sandy or loam soil, from which it would be difficult to dermally absorb PCBs. Accidental ingestion of 100 mg of this soil would result in an intake of about 1.1 ug of PCBs. An estimated average daily dietary intake of 8 to 15 ug of PCBs per day has been reported.¹⁴ Toxicity to humans from PCBs in contaminated oil was reported at an average intake of 72.4 ug per day.¹⁵ Therefore, accidental acute oral ingestion of this soil would not appear to be significant in and of itself. Volatilization of PCBs is limited by soil adsorption. Therefore, potential inhalation exposure is not expected to even approach industrial exposure levels.

An oral cancer potency of 7.7 (mg/kg/day)-1 has been developed, based on Aroclor 1260.11 Theoretical cancer risks have been calculated, based on spills covering certain measured surface areas of land and some involving inhalation exposure. Because of the nature of the PCBs (Aroclor 1248 and tentatively identified PCBs, not Aroclor 1260) and the apparent localization of contamination (one sample), it is doubtful that any useful quantitation of cancer risk could be achieved for this site. According to the no-threshold theory of carcinogenicity, however, some theoretical increase in cancer risk cannot be ruled out.

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DDT, a chlorinated hydrocarbon insecticide banned in the United States since 1973, was detected in one backyard soil sample. DDT has been seen to affect the hepatic, cardiovascular, and reproductive systems, and especially the nervous system. A dose of 20 grams has been reported to be highly dangerous but not fatal to man. 16 An oral human TDLo (lowest reported toxic dose) of 5 mg/kg has been reported. 16 The soil levels are far lower than reported toxic levels. It has been said that practically everyone born since the mid-1940s has had a lifetime exposure and storage of some DDT in their fatty tissues. 6 DDT is very lipophilic and is quite persistent in the environment. Potential DDT exposure from this soil does not appear to be significantly greater than exposure from most other sources, as DDT contamination is so widespread.

Phthalates, as plasticizers, are ubiquitous in the environment.⁶ Phthalates are noted for their low acute toxicity and low chronic toxicity.⁶ No significant impacts are expected.

1,3-Dichlorobenzene is a SVOC present at a level for which there is no evidence to suggest significant environmental impacts. It does not appear to be very persistent in the environment.

N-nitrosodiphenylamine, a moderately toxic irritant, is present at a level for which there is no evidence to suggest significant impacts.¹⁶

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